| SITE # 2832

ENGINEERING INVESTIGATIONS AT INACTIVE HAZARDOUS WASTE SITES

PHASE I INVESTIGATION

NYD 986866390

NEW YORK EMULSION TAR PRODUCTS SITE No. 633031 UTICA (C) ONEIDA (C)

DATE: FEBRUARY 1990



FILE COPY

COMPLETED

Prepared for:

NEW YORK STATE
DEPARTMENT OF ENVIRONMENTAL CONSERVATION

50 Wolf Road, Albany, New York 12233

Thomas C. Jorling, Commissioner

DIVISION OF HAZARDOUS WASTE REMEDIATION

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ENGINEERING INVESTIGATIONS AT INACTIVE HAZARDOUS WASTE SITES

IN THE STATE OF NEW YORK

PHASE I INVESTIGATION

NEW YORK EMULSION TAR PRODUCTS

CITY OF UTICA

ONEIDA COUNTY, NEW YORK

SITE No. 633031

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NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION

50 WOLF ROAD

ALBANY, NEW YORK 12233-0001

Prepared by:
URS COMPANY, INC.

570 DELAWARE AVENUE
BUFFALO, NEW YORK 14202

FEBRUARY 1990



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1.0 EXECUTIVE SUMMARY

The New York Emulsion Tar Products site is located between the Mohawk River and Utica Harbor on the north side of Utica, Oneida County, New York (Figure 1). The 2.96-acre site, presently owned by Suit-Kote Corporation, of Cortland, New York, ceased industrial operations in 1983. It was used for 57 years, first for the processing of road tars, and later for the production of asphalt emulsion. Adjacent parcels have been put to industrial uses for at least half a century (Figure 2 and Figure 3).

Investigations of the adjacent Harbor Point property by Niagara Mohawk have led to the discovery of cyanide, as well as a number of PAHs, benzene, and phenol in groundwater and soil near the site. Mohawk River sediments show signs of PAH contamination. Niagara Mohawk concluded that the New York Emulsion site appears to be one of the sources of naphthalene and benzene on the Harbor Point property, and that it may be the source of other contaminants as well.

Seven protected wetlands are found within one mile of the site. Over 100 potable water supply wells, serving an estimated 456 people, are found within a three-mile radius.

The Phase I effort involved the compilation of information gathered from several sources, including, but not limited to, the following: the New York State Department of Environmental Conservation (NYSDEC) - Region 6; the Oneida County Planning Department; and a site inspection conducted by URS Company, Inc. personnel on August 6, 1987.

The intent of the Hazard Ranking System (HRS), as developed by the Mitre Corporation under contract to the U.S. Environmental Protection Agency, is to provide a method by which uncontrolled hazardous waste sites may be systematically evaluated with regard to the potential risk

that a site may pose on human health or safety, and/or the environment. The HRS is designed to provide a numerical value through an assessment of technical data and information, and relating that information with respect to the following three hazard modes:

- o migration of hazardous substances from the site (S_M)
- o the potential for harm from fire and explosion (S_{FF})
- o the potential for harm from direct contact (S_{DC})

The migration potential (S_M) is determined through the rating of factors associated with three migration routes: groundwater (S_{gW}) , surface water (S_{SW}) and air (S_a) . The factor categories include observed release, route and waste characteristics, containment and targets. The scored value for each route is composited to determine the risk to humans and/or the environment from the migration of hazardous substances from the site (S_M) . The risks involved with the potential for fire and explosion (S_{FE}) and direct contact (S_{DC}) are evaluated according to site specific information, including: waste characteristics, containment, demographics and proximity to sensitive habitats (wetlands, wildlife, etc.).

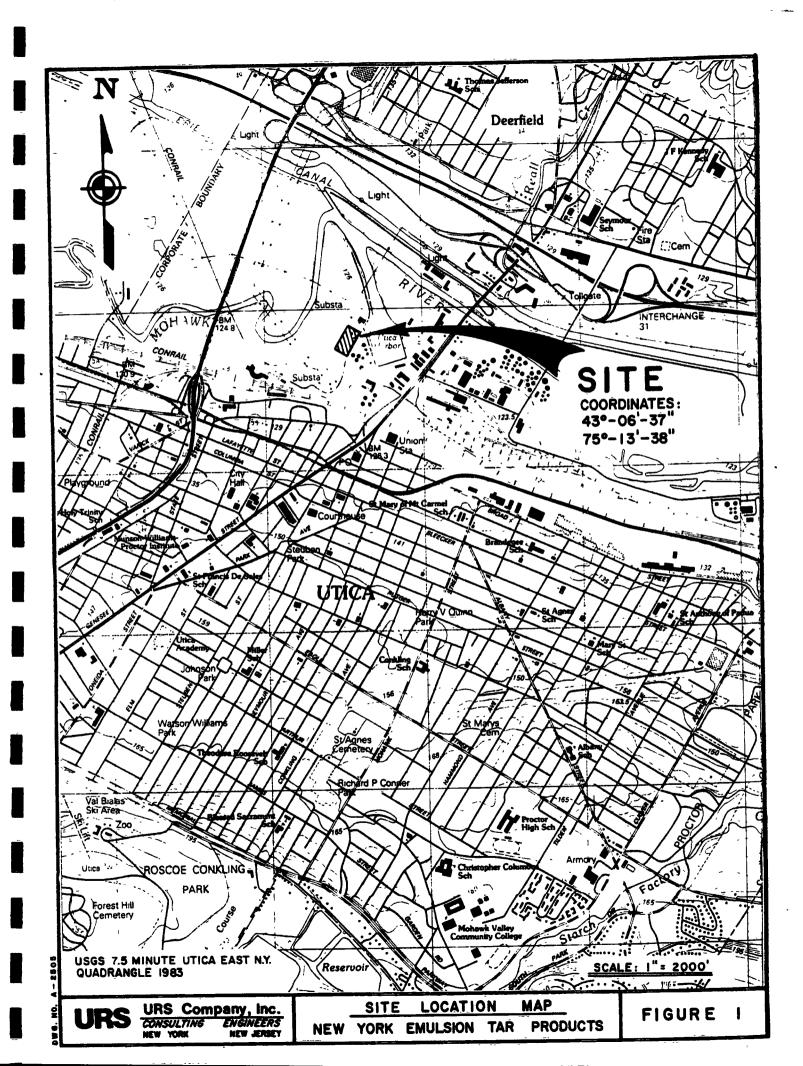
Based on information gathered during this investigation of the New York Emulsions site, the following HRS scores were obtained:

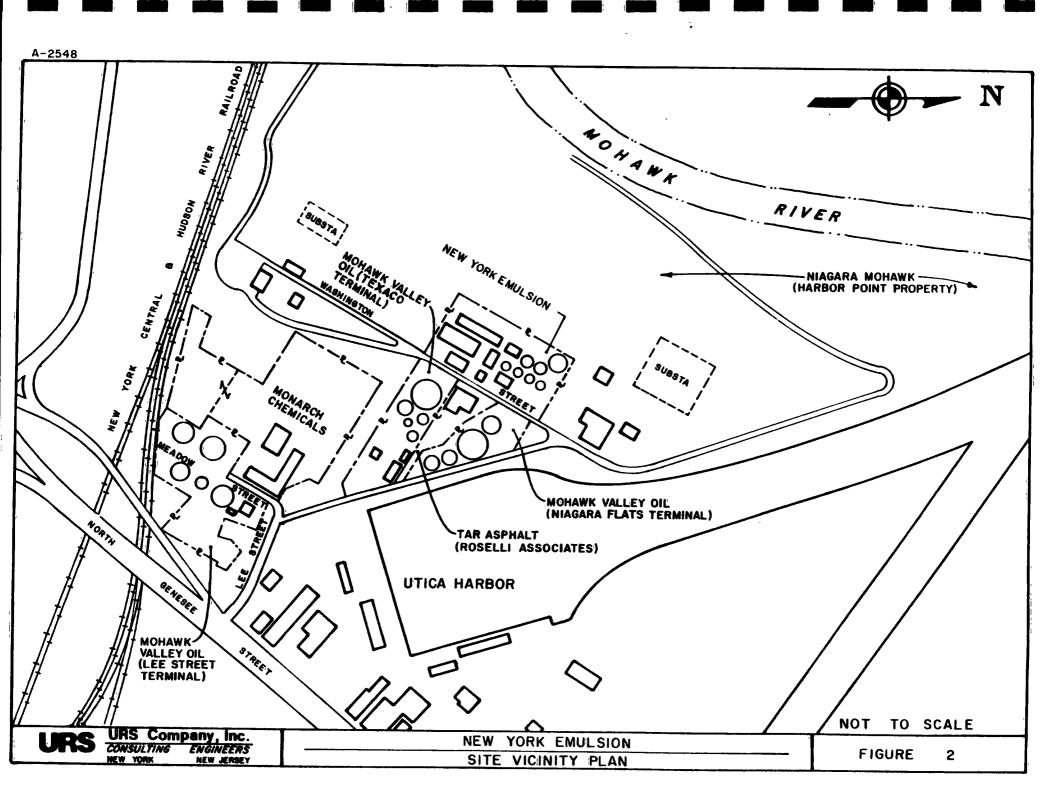
$$S_{M} = 10.74 (S_{gw} = 17.35, S_{sw} = 6.66, S_{a} = 0.00)$$

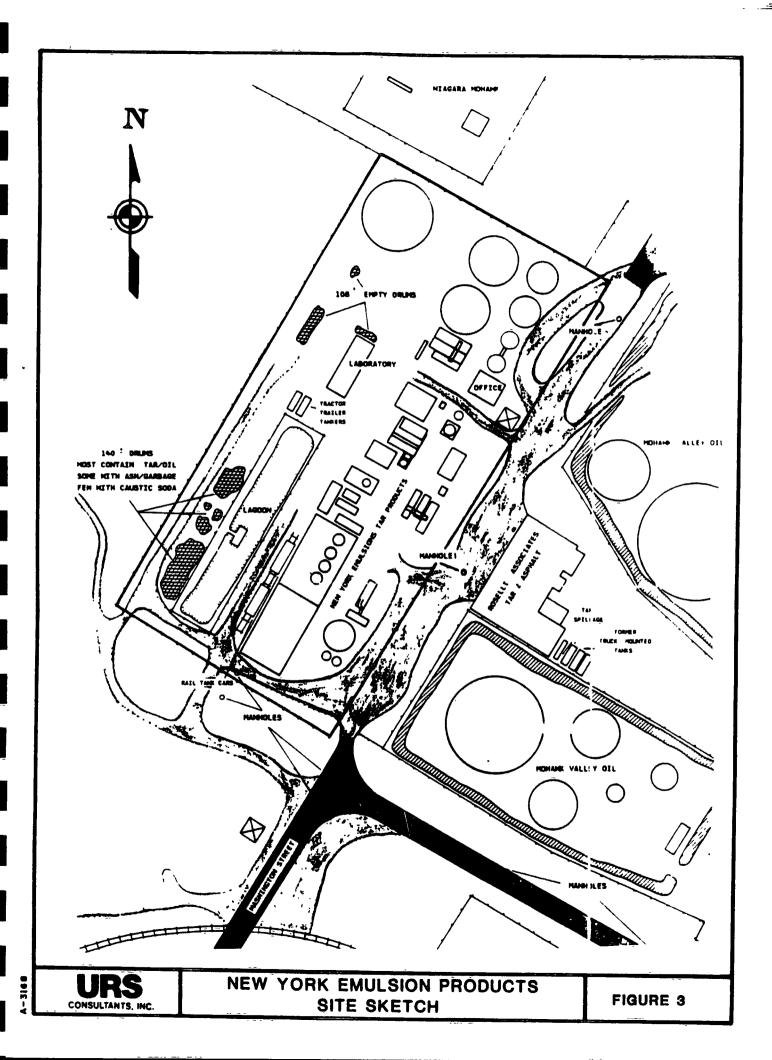
 $S_{FE} = 0.00$
 $S_{DC} = 62.50$

The data available in several areas for this Phase I investigation are considered inadequate for a proper site assessment; therefore, additional data gathering and evaluation is suggested. Proposed Phase II activities include investigation into the exact derivation of tars and oils found at the site, and subsurface investigations on site. Such

investigations should include soil borings, installation of monitoring wells, and sampling and analysis of groundwater and surface and subsurface soil.

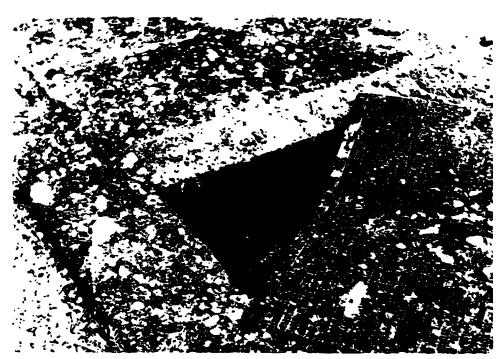








August 6, 1987 Lots of cleanup to be done here. The tanks are empty. There is some loose insulation and PCB transformers.



August 6, 1987 Lagoon discharges to manhole and storm sewer system on street. Unsure of what went into lagoon.



August 6, 1987 Hoppers used to make the material that Tar Asphalt Service would spread.



August 6, 1987 Drums on-site are mostly empty.

2.0 PURPOSE OF THE PHASE I INVESTIGATION

The Phase I investigation of the New York Emulsion Tar Products site on Washington Street was conducted for the following purposes:

- o Evaluation of the hazard and potential for harm to the public and the environment posed by the present site conditions. This includes the compilation and review of site-specific information regarding waste characteristics, routes of contaminant migration, population and/or environment at risk and operational history. All pertinent information is used to determine a Preliminary Hazard Ranking System (HRS) score for the site.
- o Evalute the adequacy of available information for the completion of a Final HRS score and identify areas where further investigation and sampling/analysis are needed to establish a valid score.

3.0 SCOPE OF WORK

The Phase I investigation at the New York Emulsion site comprised several interrelated tasks as follows:

- (a) An extensive data search was conducted, using both sitespecific and regional sources. This information was compiled from existing data, and new sources, and was used to develop a preliminary characterization of the site after review.
- (b) A site inspection was conducted in order to characterize the site and vicinity, collected information regarding the presence and disposal practices of hazardous substances (if any), photograph the site, conduct preliminary air monitoring using a Photovac TIP instrument, and confirm information obtained during the initial data search. A USEPA Site Inspection Report (EPA Form 2070-13) and the New York State Department of Environmental Conservation Inactive Hazardous Waste Disposal Site Réport were completed following the site inspection.
- (c) Preliminary HRS scores were calculated for the site and the supporting documentation records were prepared using the information obtained in the data search and site inspection.
- (d) The adequacy of available information was evaluated and recommendations were made for further investigations where necessary to properly develop a final HRS score.
- (e) The Phase I investigation report was prepared according to the terms of the contract.

During the investigation, a number of sources were contacted for information, including but not limited to:

- o Science and Engineering Library, University of Buffalo May 28, 1987, June 1, 1987 and June 24, 1987
 - Geological/hydrogeological information
- o Lockwood Library, Government Document Section, University of Buffalo June 1, 1987
 - 1980 Census information (population data and source of water data)
- o Buffalo and Erie County Public Library June 12, 1987
 - Climatalogical data and 1980 Census information
- New York State Department of Health, Syracuse Regional Office
 Ron Heerkins and Emmy Thomee June 19, 1987 and June 25, 1987
 - General files
- o New York State Museum and Science Service July 2, 1987
 - Geological maps
- o NYSDEC Watertown, Region 6 Headquarters Darrell Sweredoski July 24, 1987
 - General files
- o NYSDEC Region 6 Gregg Townsend August 4, 1987 and August 18, 1987
 - Site history and sampling data
- o NYSDEC Len Ollivett, Conservation Biologist August 6, 1987 and August 18, 1987
 - Wetlands map and endangered species information

- Oneida County Soil and Water Conservation District Robin
 Mangini August 12, 1987 and September 14, 1987
 - Irrigation and agricultural lands information
- Suit-Kote Corporation William Fowlston, Safety Director -August 18, 1987 and September 16, 1987
 - Site history
- o City of Utica Fire Chief September 2, 1987
 - Fire and explosion threat information
- o NYSDEC Albany Stuart Smith September 4, 1987
 - Site history and well location
- o Town of Frankfort Kathleen Aversa, Town Clerk September 8, 1987, September 10, 1987 and September 18, 1987
 - Source of water information
- O Utica Board of Water Supply Russell Logalbo September 10, 1987
 - Source of water information
- Town of Marcy Cindy Cochi and Karl Maxwell, Town Supervisor
 September 10, 1987 and September 15, 1987
 - Source of water information
- Oneida County Health Department Chris Demme September 10, 1987
 - Source of water information
- o Town of New Hartford Robin Rueb September 10, 1987
 - Source of water information

- o Town of Deerfield Donald S. Youlen September 10, 1987 and September 21, 1987
 - Source of water information

4.0 SITE ASSESSMENT

4.1 Site History

The 2.96-acre New York Emulsion site is located on the north side of the City of Utica, New York, between the Mohawk River and Utica Harbor. The site is adjacent to Niagara Mohawk's Harbor Point property in an area which has seen considerable industrial activity for many years (Ref. 8,9). The Niagara Mohawk property and two nearby properties, Mohawk Valley Oil and Monarch Chemicals, are currently on NYSDEC's registry of inactive hazardous waste sites. The adjacent Harbor Point property was the site of a water gas plant/coal gasification plant from 1902 to the 1950s and has undergone extensive investigations concerning the occurrence of hazardous wastes on site (Ref. 30). The Mohawk Valley Oil property was a fuel storage tank farm and is suspected as a source of napthalene and benzene contamination, while the Monarch Chemical Company is an active distribution and manufacturing plant of sodium hydroxide and sodium hypochlorite (Ref. 30).

Essentially every part of the area between the Mohawk River and Utica Harbor, in which the New York Emulsion site is located, has been disturbed at one time or another. Disturbances included construction of industrial facilities; storage of raw materials or byproducts; disposal of wastes, process byproducts, or construction materials; spills; or unauthorized dumping of materials. Prior to the industrial development of the area, the land on Harbor's Point was farmed. About 65 years ago much of the property was acquired by Utica Gas and Electric (UGE), industrial development dates back to this time. UGE acquired the land, which is currently the New York Emulsion site from the Davies family in 1923, and in 1926 sold it to American Tar Products Company (Koppers Products Company). It is not known what use UGE made of this property before selling it to American Tar Products. Utilizing the coal-tar residue of the adjacent Harbor Point coal gas production plant, Koppers

began the processing of road tar, pitches and creosote oils on site, and in later years added the production of asphalt emulsion. Tar Asphalt Service, a company located across Washington Street from the site, customarily picked up these products for use off site. Properties adjacent to or near the site have been used for storage of gasoline, kerosene, and other petroleum products; chemical manufacture; storage of coke; manufacture of water gas and coal gas; and steam generation of electricity. Large quantities of coal have in the past been stockpiled on the Harbor Point property, and railroad traffic across the area has been common. Koppers reportedly used an area just west of its property to clean road tar from its vehicles, and to dump tarry residues (Ref. 11, 12).

In March 1977 Koppers, shortly after beginning the production of emulsified asphalt, sold the New York Emulsion property and inventory to Suit-Kote Corporation, of Cortland, N.Y. Inventory at this time included both asphalt and road tar, as well as solvents including naphtha. Suit-Kote operated the asphalt emulsion plant for several years and ceased operations in 1983. Suit-Kote allegedly never processed road tar (a product of the destructive distillation of wood or coal) and confined its operations to production of asphalt emulsion (a petroleum-based product) only. Suit-Kote alleges it sold all the road tar products it had acquired with purchase of the site. The naphtha inventory was reportedly used up by Suit-Kote in the production of asphalt emulsion (Ref. 11).

Suit-Kote used only the portion of the facilities closest to Washington Street during its six years of operation (Ref. 11).

Suit-Kote alleges that no spill or accident occurred during its site tenure. The asphalt emulsion production process neither generates nor employs benzene or naphthalene, two contaminants that have been found at the Harbor Point property and attributed to the New York

Emulsion site. Suit-Kote has announced its intention of a general cleanup of the site (Ref. 11, 19).

4.2 Site Surface Characterization

The site is relatively level and is situated less than ten feet above Utica Harbor to the east and the Mohawk River to the west. The site has numerous tanks, buildings, and industrial utilities located within its boundaries and has a 0.25-acre (approximate) lagoon in the southwest quadrant of the property. The lagoon was constructed by Koppers as part of Spill Prevention, Control, and Countermeasures planning, and drains via the Lee Street storm sewer to the Mohawk River. A SPDES permit for this discharge is in effect until 1991. A number of onsite tanks have connections to a second storm sewer on Washington Street with drainage to Utica Harbor. (These two sewers were once joined.) The Washington Street sewer is apparently the only sewer draining the non-lagoon portion of the plant. Drums are located throughout the site. These drums were in rather segregated areas (near the lagoon and the lab) as shown on the site map. Most of these drums appear to be empty (Ref. 8, 11, 20, 1, and site visit).

One New York Emulsion building houses a laboratory, believed to have been used more extensively by Koppers than by Suit-Kote. Suit-Kote in fact claims that the lab was used by them for quality control only and that no exotic chemicals were employed. The laboratory does, however, contain chemicals remaining from Koppers' occupancy (Ref. 11).

Seven protected wetlands are found within one mile of the site. A Wildlife Management Area covers portions of three of these wetlands. A private wildlife preserve also exists among these wetlands. Potable water is supplied throughout nearly the entire area by the Utica Board of Water Supply, whose intakes lie 15 miles north of the site, at

Hinckley Reservoir. About twenty private wells are known in the Town of Deerfield, and 100 in the Town of Marcy. These serve an estimated 456 people (Ref. 14, 5, 6, 6a, 12).

Several utility trenches dissect the site and are in fact considered to play an important role in subsurface contaminant transport.

These trenches carry both gas and sewer lines (Ref. 1).

4.3 Site Hydrogeology

The New York Emulsion site is located within the physiographic province of the Hudson-Mohawk Lowlands. Glacial scour formed the Mohawk Valley Trough beneath the site. This was subsequently filled with a sequence of glacial till, outwash, and lake sediments. Surficial deposits in this region consist of floodplain sediments of the Mohawk River (Ref. 21, 1).

Geologic mapping indicates that bedrock of the region consists of sedimentary rocks of Middle to Upper Ordovician age. The bedrock surface underlying the site is reported as being in the Utica Shale. The Utica Shale is a finely laminated black marine shale lithology that has scattered concentrations of pyrite and a relatively high fisility. Available data indicates that groundwater occurs along bedding planes and within fractures and joints in this formation. Groundwater within the Utica Shale may contain some component of hydrogen sulfide as a result of the natural dissolution of pyrite (iron disulfide) crystals and nodules in the lithology (Ref. 22, 23).

The unconsolidated surficial deposits at the adjacent Harbor Point property start with a layer of reworked or lodgement till directly on bedrock. This layer is overlain by 90 to 120 feet of glaciolacustrine sandy silts and clay sediments. Fluvial deposits including silty sand and gravel or sandy silt overlie the lacustrine sediments and are in turn overlain by organic silt and grayish clay floodplain sediments plus

peat. The lower portions of these fluvial deposits appear to be less permeable than the upper. The entire site vicinity is covered with up to ten feet of heterogeneous fill material. Bedrock is found at depths in excess of 150 feet in areas of the Harbor Point property, but closer to the site it is found between 40 and 85 feet beneath the surface (Ref. 1, 12, 20).

Groundwater is found within all unconsolidated horizons underlying the site, although the glaciolacustrine layer appears to act to some extent as an aquiclude. Water level data from area monitoring wells indicate that the potentiometric surface is between 10 and 15 feet below the ground surface. Shallow and intermediate depth wells in the overburden (above the glaciolacustrine silt and clay) have similar water levels and are considered to be screened in the same aquifer. This overburden aquifer is believed to be isolated from the bedrock aquifer when the glaciolacustrine sediments are present, however, boring information collected near the site show that the glaciolucstrine layer is discontinuous and that there is probably connection between the bedrock and overburden aquifers. Therefore, for the purposes of HRS scoring, the overburden and bedrock aquifers will be treated collectively as the aquifer of concern (Ref. 1, 12).

Permeability within the glaciolacustrine silt and clay sediments ranges from 10^{-3} to 10^{-5} cm/sec. Due to the interaction between river stage and groundwater level on the Harbor Point property, it is difficult to estimate the average rate and direction of groundwater flow. Groundwater movement has at times been observed from the river towards the center of the property. Net movement nevertheless appears to be toward the river, at a very low rate (Ref. 1).

4.4 Site Contamination

Heavy tar-product accumulations were noted in sediments dredged from Utica Harbor by the New York State Department of Transportation in

April 1981. Naphthalene, acenaphthalene, phenanthrene, and anthracene were identified in this tar substance. These constituents are not characteristic of "oil-derived" tars and therefore it is unlikely the tar material was produced in the asphalt emulsion plant. Site assessments were conducted on the Harbor Point property and surrounding areas (including the NY Emulsions site) in December 1983 by Niagara Mohawk and in January 1984 by USEPA to identify the probably source of the contamination. Based on results of these assessments, Niagara Mohawk commissioned a site investigation and produced the first in a series of reports in March 1984. The investigation covered contamination of both land and river. In addition, NYSDEC sampled tars at and near the NY Emulsion site in 1985, demonstrating exactness of match between tars found on and nearby off site. A naphthalene peak was isolated in both onsite and offsite samples (Ref. 18, 12, 24).

Subsurface investigations by Niagara Mohawk on the Harbor Point property revealed a surficial layer of waste material (3-6 feet deep in the area of the New York Emulsion site) consisting of building debris, gas production process wastes (i.e., coal-tar related materials), and other miscellaneous wastes including municipal landfill refuse. Wastes in the center of the Harbor Point property and west of the New York Emulsion site, contained relatively high levels of polynuclear aromatics (PAHs). Soil concentrations of naphthalene reached 34,300 mg/kg near the southwest corner of the New York Emulsion site in the area where Koppers allegedly cleaned vehicles and deposited road tar materials. A hard "asphalt-like" substance was observed at or near the ground surface here. Oils and tars were evident in test pits and monitoring wells on the periphery of the New York Emulsion site (Ref. 20, 1).

Groundwater studies showed highest Total Organic Carbon levels near the New York Emulsion site and also near a gas storage holder 375 feet west of the plant on the Harbor Point property. Test pits showed tar to be concentrated at or near the soil surface. Contamination in the soil was more widespread in the upper fill layers than in the underlying fluvial silts and clays while contamination in groundwater was confined to the shallow and intermediate levels. Most contaminants are thought to be confined to the fill and alluvial horizons and not in the bedrock. Niagara Mohawk constructed isoconcentration plots from groundwater data that illustrated the centers of phenol, naphthalene, acenaphthylene, and acenaphthene concentration downgradient of the New York Emulsion site. All these substances are coal-tar derivatives. The Niagara Mohawk first-round report pointed to a "common source" of acenaphthene and acenaphthylene "in the center of the site between the emulsions plant and the gas storage holder" while the second-round report judged there to be two sources of these materials (Ref. 20, 1, 12).

No organic vapors were detected on site during 1984 investigations (Ref. 12).

Investigations of the Mohawk River could not demonstrate that water quality had been solely affected by the Harbor Point property. Sediments near the point contained measurable levels of PAHs, but the picture was complicated by the discovery of acenaphthylene in sediments 1,000 feet upstream as well. Major PAH sediment concentrations were found in Utica Harbor and at the discharge of the Lee Street sewer into the river (Ref. 25).

On the basis of first-round findings, the entire northern part of the Harbor Point property was eliminated from further consideration, and the investigative effort was brought to bear more directly upon the area surrounding the New York Emulsion property, the gas storage tank, and sewer lines. It was stated that first-round investigations had indicated the possibility of a "petroleum-based source" of contamination (presumably a reference to the emulsified asphalt operation at the New York Emulsion site), and investigators therefore announced plans for a special effort to determine the makeup of tars and oils found at these

locations--i.e to determine whether they were coal-tar based as had originally been thought, or petroleum-based. Results of investigations on this critical point, however, were not available during this Phase I investigation (Ref. 1).

Twenty-nine monitoring wells were constructed and sampled for volatiles and other compounds in conjunction with the second-round investigation at the Harbor Point property. Cyanide was definitively attributed to the former gas purifier boxes west of the NY Emulsion site, on Niagara Mohawk property; naphthalene was, on the basis of groundwater data, attributed by Niagara Mohawk to both the New York Emulsion site and an offsite source to the north; New York Emulsion appeared to investigators as a source of benzene which was measured in groundwater just east of the NY Emulsion site at 11,000 ppb. Contaminant loading to the river was determined to be heaviest from the sewers, with transport by groundwater being of secondary importance. The contribution from surface runoff was deemed insignificant (Ref. 1, 2).

Investigations of the adjacent Harbor Point property by Niagara Mohawk, as well as a surficial tar sample collected on the New York Emulsion site by NYSDEC in 1985, indicate that at least a portion of the observed (benzene and naphthalene) groundwater contamination at Harbor Point is attributable to New York Émulsion. Because the production of road tars (by Koppers Products Company) at the New York Emulsion site involved utilization of the coal tar byproduct from the Harbor Point coal gasification plant, it is difficult to separate these two sites from one another for the purpose of establishing the source of coal tar-related contaminants in area groundwater. However, it does not appear likely that the occurrence of benzene and naphthalene in Harbor Point groundwater samples is related to the more recent production of emulsified asphalt at the New York Emulsion site, since the emulsified asphalt process reportedly neither uses nor produces either compound. While both naphthalene and benzene may be petroleum-derived (as is

asphalt itself) or coal tar-derived, they are isolated from petroleum only after catalytic reprocessing of the lighter fractions. Again, however, regardless of the time of occurrence or process involved at that time, the benzene and naphthalene contamination of area groundwater is considered to be at least partially attributable to the New York Emulsion site. On this basis, an HRS score for the site has been developed using benzene and naphthalene as observed contaminant releases to groundwater (Ref. 11, 26).

NARRATIVE

5.0 PRELIMINARY APPLICATION OF THE HAZARD RANKING SYSTEM

5.1 Narrative Summary

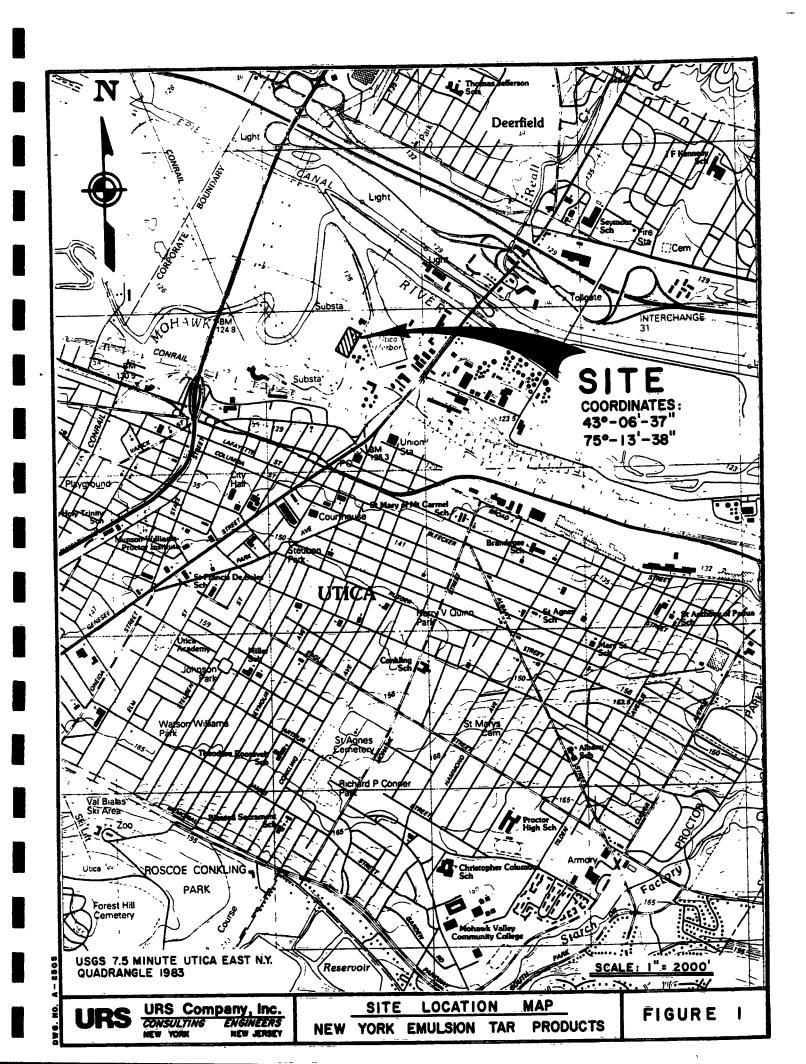
New York Emulsion Tar Products Washington Street Utica (C) Oneida (C) New York

The 2.96-acre New York Emulsion Tar Products site is located on the north side of the City of Utica, New York, between the Mohawk River and Utica Harbor. It is presently owned by Suit-Kote Corporation, which processed emulsified asphalt there from 1977 to 1983. The site's previous owner, Koppers Products Company, processed both road tar products and asphalt emulsion prior to turning it over to Suit-Kote.

No waste products were associated with either operation, although hazardous materials attributable to tar products production have been found in soils and groundwater near and around the site. A number of related materials have also been found in Mohawk River sediments. These substances have included PAHs and volatile organics. The site is located in an area where such materials were used or produced at various facilities on adjoining properties. Protected wetlands are found within one mile of the site, and over 100 potable water wells within three miles.

Niagara Mohawk, owns one parcel of adjacent land known as Harbor Point and has been investigating contamination on that property since early 1984. Some of their investigations extend onto the New York Emulsion property.

LOCATION



Facility name: New York Emulsion Tar Products
Location: Washington Street, Utica (c), Oneida County
EPA Region:
Person(s) in charge of the facility: William Fowlston, Director of Safety & Environment
Name of Reviewer: URS Corporation Date: November 11, 1987
General description of the facility: (For example: landfill, surface impoundment, pile, container; types of hazardous substances; location of the facility; contamination route of major concern; types of information needed for rating; agency action, etc.)
Originally site of manufacture and distribution of tar products,
and later of asphalt emulsion. Investigations around perimeter of
site have led to conclusion that site is source of naphthalene
and benzene. Transport of contaminants to nearby surface water
via storm drains and groundwater. Further investigation required
on site itself .
•
Scores: S _M = 10.74 17.35 6.66 0.00
SFE = 0.00
Spc = 62.50

HRS COVER SHEET

			Ground Wat	er Route Wo	rk Shee				and the second of the second o
	Rating Factor	,	Assigned Value (Circle One)				Score	Max. Score	Ref. (Section)
1	Observed Release		0 45				45	45	3.1
	If observed releas	-		•					
2	Route Characteris Depth to Aquifer Concern		0 1 2	3	7	2		6	3.2
	Net Precipitation Permeability of t Ungaturated Zo	the	0 1 2 0 1 2	3		1		3	
	Physical State		0 1 2			1	-	3	
	4	MATERIAL SERVICES	Total Route Ch	eracteristics	Score			15	
3	Containment		0 1 2	3		1	<u> </u>	3	3.3
4	Waste Characteria Toxicity/Persiste Hezardous Wast Quantity	ence	0 3 6	9 12 15 18 3 4 5 6	7 8	1	1 <u>2</u> 1	18	3.4
	· · · · · · · · · · · · · · · · · · ·		Total Waste Ch	aracteristics	Score		13	26	
3	Targets Ground Water U: Distance to Near Well / Population Served	rest	12 16 1	2 3 6 8 10 18 29 12 35 40		3	9	9 40	3.5
<u></u>	If line 🛐 is 45.	multiply	Total Ta	rgets Score			17	49	
	If line 1 is 0, m						9,945	57,330	
7	Divide line 6 by	y 57,330 a	nd multiply by	100		Sgw-	17.3	5	

GROUND WATER ROUTE WORK SHEET

	-	Surface Water Route Work She	et	. 90 4 2		
	Rating Factor	· Assigned Value (Circle One)	Multi- plier	Score	Max. Score	Ref. (Section
1	Observed Release	0 45	1	0	46	4.1
		is given a value of 45, proceed to line 4: is given a value of 0, proceed to line 2.		-		
2	Route Characteristic Facility Slope and	~	1	0	3	4.2
	Terrain 1-yr. 24-hr. Rainfa Distance to Neare		1 2	2 6	3	
	Water Physical State	0 1 2 3	1	3	\$	
		Total Route Characteristics Score		11	16	
3	Containment	0 1 2 3	1	3	3	4.3
U	Waste Characteristic Toxicity/Persister Hazardous Waste Quantity	0 3 6 9(12)15 18	1	12	18	4.4
		Total Waste Characteristics Score		13	26	
3)	Targets Surface Water Use Distance to a Sens		3 2 -	6 4	9	4.5
	Population Served to Water Intake Downstream	7/Distance 0 4 6 8 10 12 16 18 20 24 30 32 35 40	1	0	40	
		Total Targete Score		10	55	
		uitiply [] x 4 x 5 itiply 2 x 3 x 4 x 5		4290	64,350	•
7]	Divide line 6 by	64,350 and multiply by 100	S _{SW} -			<u></u>

SURFACE WATER ROUTE WORK SHEET

•	Air Route Work	k Sheet	_		
Rating Factor	Assigned Value (Circle One)	Mult plie	I 600	Max. Score	Ref. (Section
Observed Release	0 48	1	Ô	45	5.1
Date and Location:					
Sampling Protocol:			_		
If line $\boxed{1}$ is 0, the \overline{S}_{a} if line $\boxed{1}$ is 45, then	= 0. Enter on line 3. proceed to line 2.				
Waste Characteristics					5.2
Reactivity and Incompatibility	0 1 2 3	1		3	
Toxicity Hazardous Waste Quantity	0 1 2 3 0 1 2 3 4 (3 6 7 8 1		9	•
	Total Waste Characterist	cs Score		T 1	-
3 Targets			<u> </u>	20	<u> </u>
Population Within	į 0 9 12 15 18	1		30	5.3
4-Mile Radius Distance to Sensitive	21 24 27 30 0 1 2 .3	•		30	
Environment Land Use		2		6	
Lang USS	0 1 2 3	1		3	
	Total Targets Score			39	
Multiply 1 x 2 x 3	3	200	→	25 100	
	_			35,100	
Divide line 4 by 35,100	and multiply by 100	Sa-	0.00		

AIR ROUTE WORK SHEET

	8	s ²
Groundwater Route Score (Sgw)	17.35	301.02
Surface Water Route Score (S _{SW})	6.66	43.36
Air Route Score (Sa)	0.00	0.00
$s_{gw}^2 + s_{sw}^2 + s_a^2$		345.38
$\sqrt{s_{gw}^2 + s_{sw}^2 + s_a^2}$		18.58
$\sqrt{s_{gw}^2 + s_{sw}^2 + s_a^2} / 1.73 - s_M -$		10.74

WORKSHEET FOR COMPUTING SM

NO DOÇUMENTED FIRE OR EXPLOSION THREAT

Rating Factor		Ass (C		ed (Multi- plier	Score	Max. Score	Ref. (Section
Containment	1					3		1	•	3	7.1
Waste Characteristics Direct Evidence ignitability Reactivity incompetibility Hazardous Waste Quantity	0	1 1	2	3	4	5 (. 7 8	1 1 1 1		3 3 3 3 8	7.2
	Total Wa	ște -	Cha	rac	teri	stics	Score		_	20	
Targets Distance to Nearest Population		1			4	5		1		5	7.3
Distance to Nearest Building Distance to Sensitive		1		_				1		3	
Environment Land Use Population Within 2-Mile Radius Buildings Within	. 0	1 1	2	3		•		1 1		3 5	
2-Mile Radiuș										·	
Multiple (i)	Tot	al T	arg	ets	Sci	070	· <u>-</u>			24	
Multiply 1 x 2 x 3 Divide line 4 by 1,440 and									<u> </u>	1,440	

FIRE AND EXPLOSION WORK SHEET

		Direct Contact Work She	et			
	Rating Factor	Assigned Value (Circle One)	Multi- plier	Score	Max. Score	Ref. (Section)
1	Observed Incident	0 45	1	0	45	8.1
	If line 1 is 45, proceed to	_4,				
2	Accessibility	0 1 2 3	1	3	3	8.2
3	Containment	0 (15)	1	15	15	8.3
4	Waste Characteristics Toxicity	0 1 2(3)	5	15	15	8.4
5	Targets Population Within a 1-Mile Radius	0 1 2 3 4 8	4	20	20	8.5
	Distance to a Critical Habitat	0123	4	0	12	
	·	U				
		Total Targets Score	1.	20	32	
_	line 1 is 45, multiply [Total Targets Score		20 500 2	32	

DIRECT CONTACT WORK SHEET

5.4 HRS Documentation Records

FACILITY NAME:	New York Emulsion Tar Products	
LOCATION:	Washington Street, Utica	
	Oneida County, New York	

GROUNDWATER ROUTE

- 1 OBSERVED RELEASE
- o CONTAMINANTS DETECTED (5 MAXIMUM):

Naphthalene, benzene (Ref. 1, 2).

RATIONALE FOR ATTRIBUTING THE CONTAMINANTS TO THE FACILITY:

Contaminants measured in shallow and medium depth groundwater in vicinity of site. Isoconcentration plot lines elevated around site perimeter (Ref. 1, 2).

Score 45

2 ROUTE CHARACTERISTICS

DEPTH TO AQUIFER OF CONCERN

o NAME/DESCRIPTION OF AQUIFER(S) OF CONCERN:

Observed release precludes evaluation of this factor.

O DEPTH(S) FROM THE GROUND SURFACE TO THE HIGHEST SEASONAL LEVEL OF THE SATURATED ZONE [WATER TABLE(S)] OF THE AQUIFER OF CONCERN:

Observed release precludes evaluation of this factor.

O DEPTH FROM THE GROUND SURFACE TO THE LOWEST POINT OF WASTE DIS-POSAL/STORAGE:

Observed release precludes evaluation of this factor.

NET PRECIPITATION

- o MEAN ANNUAL OR SEASONAL PRECIPITATION (LIST MONTHS FOR SEASONAL):
 Observed release precludes evaluation of this factor.
- MEAN ANNUAL LAKE OR SEASONAL EVAPORATION (LIST MONTHS FOR SEASON-AL):

Observed release precludes evaluation of this factor.

o NET PRECIPITATION (SUBTRACT THE ABOVE FIGURES):

Observed release precludes evaluation of this factor.

Score 0

PERMEABILITY OF UNSATURATED ZONE

- o SOIL TYPE IN UNSATURATED ZONE:

 Observed release precludes evaluation of this factor.
- o PERMEABILITY ASSOCIATED WITH SOIL TYPE:

 Observed release precludes evaluation of this factor.

Score 0

PHYSICAL STATE

O PHYSICAL STATE OF SUBSTANCES AT TIME OF DISPOSAL (OR AT PRESENT TIME FOR GENERATED GASES):

Observed release precludes evaluation of this factor.

3 CONTAINMENT

CONTAINMENT

METHOD(S) OF WASTE OR LEACHATE CONTAINMENT EVALUATED:

Observed release precludes evaluation of this factor.

o METHOD WITH HIGHEST SCORE:

Observed release precludes evaluation of this factor.

Score 0

4 WASTE CHARACTERISTICS

TOXICITY AND PERSISTENCE

o COMPOUND(S) EVALUATED:

Naphthalene and benzene (Ref. 1, 2).

O COMPOUND WITH HIGHEST SCORE:

Benzene

	Toxicity	Persistence	Score
Benzene	3	1	12
Naphthaler	e 2	1	9

(Ref. 3, 4)

Score 12

HAZARDOUS WASTE QUANTITY

O TOTAL QUANTITY OF HAZARDOUS SUBSTANCES AT THE FACILITY, EXCLUDING THOSE WITH A CONTAINMENT SCORE OF O (GIVE A REASONABLE ESTIMATE EVEN IF QUANTITY IS ABOVE MAXIMUM):

Unknown. Minimum non-zero amount presumed.

Score 1

O BASIS OF ESTIMATING AND/OR COMPUTING WASTE QUANTITY:

N/A

5 TARGETS

GROUNDWATER USE

o USE(S) OF AQUIFER(S) OF CONCERN WITHIN A 3-MILE RADIUS OF THE FACILITY:

Drinking water. No municipal water from alternate unthreatened sources presently available (Ref. 5).

Score 3

DISTANCE TO NEAREST WELL

 LOCATION OF NEAREST WELL DRAWING FROM <u>AQUIFER OF CONCERN</u> OR OC-CUPIED BUILDING NOT SERVED BY A PUBLIC WATER SUPPLY:

Nearest known well on Smith Hill Road, Town of Deerfield (Ref. 5, 6a).

o DISTANCE TO ABOVE WELL OR BUILDING:

Approximately 2.8 miles (Distance Value = 1) (Ref. 5).

POPULATION SERVED BY GROUNDWATER WELLS WITHIN A 3-MILE RADIUS

o IDENTIFIED WATER-SUPPLY WELL(S) DRAWING FROM AQUIFER(S) OF CONCERN WITHIN A 3-MILE RADIUS AND POPULATIONS SERVED BY EACH:

Smith Hill Road and Walker Road, Town of Deerfield; Town of Marcy (Ref. 5, 6a).

COMPUTATION OF LAND AREA IRRIGATED BY SUPPLY WELL(S) DRAWING FROM AQUIFER(S) OF CONCERN WITHIN A 3-MILE RADIUS, AND CONVERSION TO POPULATION (1.5 PEOPLE PER ACRÉ):

None (Ref. 7).

o TOTAL POPULATION SÉRVED BY GROUNDWATER WITHIN A 3-MILE RADIUS:

Town of Deerfield, 76; Town of Marcy, 380. Total 456. (Population value = 2) (Ref. 3, 5, 6a)

SURFACE WATER ROUTE

- 1 OBSERVED RÉLEASE
- O CONTAMINANTS DETECTED IN SURFACE WATER AT THE FACILITY OR DOWNHILL FROM IT (5 MAXIMUM):

None observed.

O RATIONALE FOR ATTRIBUTING THE CONTAMINANTS TO THE FACILITY:

N/A

Score 0

2 ROUTE CHARACTERISTICS

FACILITY SLOPE AND INTERVENING TERRAIN

O AVERAGE SLOPE OF FACILITY IN PERCENT:

Facility relatively level (Ref. site visit).

NAME/DESCRIPTION OF NEAREST DOWNSLOPE SURFACE WATER:

Utica Harbor is the nearest downslope surface water body and is located east of the NY Emulsions facility. The harbor is maintained by the Department of Transportation and is connected to the Mohawk River (Ref. 8, 9, 10.) In addition, storm sewers around the site discharge to the harbor.

• AVERAGE SLOPE OF TERRAIN BETWEEN FACILITY AND ABOVE-CITED SURFACE WATER BODY IN PERCENT:

Approximately 1% slope variable to north, northwest and northeast (Ref. 8).

O IS THE FACILITY LOCATED EITHER TOTALLY OR PARTIALLY IN SURFACE WATER?

No (Ref. 8 and site visit).

O IS THE FACILITY COMPLETELY SURROUNDED BY AREAS OF HIGHER ELEVATION? No, the area has low relief, gently sloping variably to the east, west, and northwest (Ref. 8).

1-YEAR 24-HOUR RAINFALL IN INCHES

2.3 inches (Ref. 3).

Score 2

DISTANCE TO NEAREST DOWNSLOPE SURFACE WATER

Utica harbor is approximately 250 feet east of the site (Ref. 8).

Score 3

PHYSICAL STATE OF WASTE

Viscous liquid (Ref. 11, 12).

Score 3

3 CONTAINMENT

CONTAINMENT

- o METHOD(S) OF WASTE OR LEACHATE CONTAINMENT EVALUATED: Landfill only (Ref. site visit).
- o MÉTHOD WITH HIGHEST SCORE:

Landfill. Not covered, no diversion system present (Ref. 3).

Score 3

4 WASTE CHARACTERISTICS

TOXICITY AND PERSISTENCE

o COMPOUND(S) EVALUATED

Naphthalene and benzene (Ref. 1, 2).

o COMPOUND WITH HIGHEST SCORE:

Benzene

Benzene

Benzene

Naphthalene

Toxicity Persistence Score
1 12
Naphthalene
2 1 9

(Ref. 3, 4)

Score 12

HAZARDOUS WASTE QUANTITY

O TOTAL QUANTITY OF HAZARDOUS SUBSTANCES AT THE FACILITY EXCLUDING THOSE WITH A CONTAINMENT SCORE OF O (GIVE A REASONABLE ESTIMATE EVEN IF QUANTITY IS ABOVE MAXIMUM):

Unknown. Minimum non-zero amount presumed.

Score 1

o BASIS OF ESTIMATING AND/OR COMPUTING WASTE QUANTITY:

N/A

5 TARGETS

SURFACE WATER USE

O USE(S) OF SURFACE WATER WITHIN 3 MILES DOWNSTREAM OF THE HAZARDOUS SUBSTANCE:

Fishing and boating (Ref. 13, 3).

Score 2

o IS THERE TIDAL INFLUENCE?

There is no tidal influence on the site (Ref. 8).

DISTANCE TO A SENSITIVE ENVIRONMENT

DISTANCE TO 5-ACRE (MINIMUM) COASTAL WETLAND, IF 2 MILES OR LESS:

None within 2 miles of the site (Ref. 14).

O DISTANCE TO 5-ACRE (MINIMUM) FRESH-WATER WETLAND, IF 1 MILE OR LESS:

Fresh-water wetlands UE-2, UE-3, UE-4, UE-5, UE-6, UE-9, and UE-10 are within one mile of the site. Fresh-water wetland UE-9 is the closest at 1,200 feet northwest of the site (Ref. 14).

O DISTANCE TO CRITICAL HABITAT OF AN ENDANGERED SPECIES OR NATIONAL WILDLIFE REFUGE, IF 1 MILE OR LESS:

None within 1 mile (Ref. 14).

Score 2

POPULATION SÉRVÉD BY SURFACE WATER

O LOCATION(S) OF WATER-SUPPLY INTAKE(S) WITHIN 3 MILES (FREE-FLOWING BODIES) OR 1 MILE (STATIC WATER BODIES) DOWNSTREAM OF THE HAZARDOUS SUBSTANCE AND POPULATION SERVED BY EACH INTAKE:

No water supply intakes within 3 miles of the site (Ref. 15, 12).

O COMPUTATION OF LAND AREA IRRIGATED BY ABOVE-CITED INTAKE(S) AND CONVERSION TO POPULATION (1.5 PEOPLE PER ACRE):

N/A

o TOTAL POPULATION SERVED:

N/A

O NAME/DESCRIPTION OF NEAREST OF ABOVE WATER BODIES:

N/A

o DISTANCE TO ABOVE-CITED INTAKES, MEASURED IN STREAM MILES.

N/A

AIR ROUTE

- 1 OBSERVED RELEASE
- O CONTAMINANTS DETECTED:

Limited air monitoring data available. Photovac TIP measurements for organic vapors taken during site inspection. None detected (\mathring{R} ef. site visit).

DATE AND LOCATION OF DETECTION OF CONTAMINANTS

No observed air release.

o METHODS USED TO DETECT THE CONTAMINANTS:

No observed air release.

o RATIONALE FOR ATTRIBUTING THE CONTAMINANTS TO THE SITE:

No observed air release.

Score 0

2 WASTE CHARACTERISTICS

REACTIVITY AND INCOMPATIBILITY

o MOST REACTIVE COMPOUND:

No observed air release.

o MOST INCOMPATIBLE PAIR OF COMPOUNDS:

No observed air release.

TOXICITY

o MOST TOXIC COMPOUND:

No observed air release.

Score 0

HAZARDOUS WASTE QUANTITY

o TOTAL QUANTITY OF HAZARDOUS WASTE:

No observed air release.

Score 0

BASIS OF ESTIMATING AND/OR COMPUTING WASTE QUANTITY:

No observed air release.

3 TARGETS

POPULATION WITHIN 4-MILE RADIUS

O UNDERLINE RADIUS USED, GIVE POPULATION, AND INDICATE HOW DETERMINED:

0 to 4 mi 0 to 1 mi 0 to 1/2 mi 0 to 1/4 mi

No observed air release.

Score 0

DISTANCE TO A SENSITIVE ENVIRONMENT

o DISTANCE TO 5-ACRE (MINIMUM) COASTAL WETLAND, IF 2 MILES OR LESS: No observed air release. o DISTANCE TO 5-ACRE (MINIMUM) FRESH-WATER WETLAND, IF 1 MILE OR LESS:

No observed air release.

O DISTANCE TO CRITICAL HABITAT OF AN ENDANGERED SPECIES, IF 1 MILE OR LESS:

No observed air release.

Score 0

LAND USE

o DISTANCE TO COMMERCIAL/INDUSTRIAL AREA, IF 1 MILE OR LESS:

No observed air release.

O DISTANCE TO NATIONAL OR STATE PARK, FOREST, OR WILDLIFE RESERVE, IF 2 MILES OR LESS:

No observed air release.

O DISTANCE TO RESIDENTIAL AREA, IF 2 MILES OR LESS:

No observed air release.

O DISTANCE TO AGRICULTURAL LAND IN PRODUCTION WITHIN PAST 5 YEARS, IF 1 MILE OR LESS:

No observed air release.

O DISTANCE TO PRIME AGRICULTURAL LAND IN PRODUCTION WITHIN PAST 5 YEARS, IF 2 MILES OR LESS:

No observed air release.

O IS A HISTORIC OR LANDMARK SITE (NATIONAL REGISTER OR HISTORIC PLACES AND NATIONAL NATURAL LANDMARKS) WITHIN THE VIEW OF THE SITE?

No observed air release.

FIRE AND EXPLOSION

- 1 CONTAINMENT
- O HAZARDOUS SUBSTANCES PRESENT:

No documented fire or explosion threat (Ref. 16).

O TYPE OF CONTAINMENT, IF APPLICABLE

No documented fire or explosion threat.

Score 0

2 WASTE CHARACTERISTICS

DIRECT EVIDENCE

TYPE OF INSTRUMENT AND MEASUREMENTS:

No documented fire or explosion threat.

Score 0

IGNITABILITY

o COMPOUND USED:

No documented fire or explosion threat.

Score 0

REACTIVITY

o MOST REACTIVE COMPOUND:

No documented fire or explosion threat.

Score 0

INCOMPATIBILITY

o MOST INCOMPATIBLE PAIR OF COMPOUNDS:

No documented fire or explosion threat.

HAZARDOUS WASTE QUANTITY

o TOTAL QUANTITY OF HAZARDOUS SUBSTANCES AT THE FACILITY:
No documented fire or explosion threat.

Score 0

o BASIS OF ESTIMATING AND/OR COMPUTING WASTE QUANTITY:
No documented fire or explosion threat.

3 TARGETS

DISTANCE TO NEAREST POPULATION

No documented fire or explosion threat.

Score 0

DISTANCE TO NEAREST BUILDING

No documented fire or explosion threat.

Score 0

DISTANCE TO SENSITIVE ENVIRONMENT

DISTANCE TO WETLANDS:

No documented fire or explosion threat.

O DISTANCE TO CRITICAL HABITAT:

No documented fire or explosion threat.

LAND USE

DISTANCE TO COMMERCIAL/INDUSTRIAL AREA, IF 1 MILE OR LESS:

No documented fire or explosion threat.

O DISTANCE TO NATIONAL OR STATE PARK, FOREST, OR WILDLIFE RESERVE, IF 2 MILES OR LESS:

No documented fire or explosion threat.

o DISTANCE TO RESIDENTIAL AREA, IF 2 MILES OR LESS:

No documented fire or explosion threat.

O DISTANCE TO AGRICULTURAL LAND IN PRODUCTION WITHIN PAST 5 YEARS, IF 1 MILE OR LESS:

No documented fire or explosion threat.

O DISTANCE TO PRIME AGRICULTURAL LAND IN PRODUCTION WITHIN PAST 5 YEARS, IF 2 MILES OR LESS:

No documented fire or explosion threat.

O IS A HISTORIC OR LANDMARK SITE (NATIONAL REGISTER OR HISTORIC PLACES AND NATIONAL NATURAL LANDMARKS) WITHIN THE VIEW OF THE SITE?

No documented fire or explosion threat.

Score 0

POPULATION WITHIN 2-MILE RADIUS

No documented fire or explosion threat.

Score 0

BUILDINGS WITHIN 2-MILE RADIUS

No documented fire or explosion threat.

DIRECT CONTACT

1 **OBSERVED INCIDENT** DATE, LOCATION, AND PERTINENT DETAILS OF INCIDENT: None observed. Score 0 2 **ACCESSIBILITY** DESCRIBE TYPE OF BARRIER(S): Barrier does not completely surround facility (Ref. site visit). Score 3 *** 3 CONTAINMENT TYPE OF CONTAINMENT, IF APPLICABLE: Materials have been spilled on ground (Ref. 12). Score 15 WASTE CHARACTERISTICS TOXICITY COMPOUNDS EVALUATED: Naphthalene and benzeně (Řef. 1, 2). COMPOUND WITH HIGHEST SCORE: Benzene Toxicity Benzene Naphthalene (Ref. 3, 4)

5 TARGETS

POPULATION WITHIN ONE-MILE RADIUS

Estimated at 18,908 (Ref. 8, 17).

Score 5

DISTANCE TO CRITICAL HABITAT (OF ENDANGERED SPECIES)

None within 1 mile of the site (Ref. 14).

Score $\underline{\underline{0}}$

DATA SOURCES AND REFERENCES

- 1. Harbor Point Property Land Investigations, Results of Extended Site Investigations, Step 3 Land Report, Niagara Mohawk Power Corporation, March 1985 (Revised May 1985).
- 2. Harbor Point Property River Investigations, Remedial Design Concepts, Step 4 River Report, Niagara Mohawk Power Corporation, June 1986.
- 3. Uncontrolled Hazardous Waste Site Ranking System, A Users Manual (HW-10), United States Environmental Protection Agency 1984, Figures 4 and 8, Tables 2, 3, 4, 5, and 9, pp. 24, 25, and 27.
- 4. Dangerous Properties of Industrial Materials Sixth Edition, N. Irving Sax, Van Nostrand Reinhold Company, New York, 1984.
- 5. Map and Statement by Donald S. Youlen, Supervisor, Town of Deerfield, New York, October 26, 1987.
- 6. Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, to Russell Logalbo, Utica Board of Water Supply, September 10, 1987.
- 5a. Karl P. Maxwell, Supervisor, Town of Marcy, New York, to Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, November 12, 1987.
- 7. Robin Mangini, District Conservationist, United States Department of Agriculture, Soil Conservation Service, to Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, September 14, 1987.
- 8. USGS Topographic Maps, 7.5 Series: Utica East, New York, Quadrangle, 1983; Utica West, New York, Quadrangle, 1955; South Trenton, New York, Quadrangle, 1983; Oriskany, New York, Quadrangle, 1955.

- 9. City of Utica, Oneida County, New York Tax Map, Sheet No. 318.08.
- 10. City of Utica, Oneida County, New York Aerial Maps, Sheet Nos. 306, 318.
- 11. William Fowlston, Safety Director, Suit-Kote Corporation, to Daniel W. Rothman, Phase I Investigation Project Manager, URS Corporation, September 16, 1987.
- 12. Harbor Point Property Land Investigations, Proposal for Initial Site Survey, Step 1 Land Report, Niagara Mohawk Power Corporation, March 1984.
- 13. Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, telecon to James Doyle, Sanitary Engineer, NYSDEC Region 6, October 8, 1987.
- Leonard E. Ollivett, Conservation Biologist, NYSDEC Region 6, to Linda J. Clark, Project Geologist, URS Corporation, August 18, 1987.
- 15. New York State Atlas of Community Water System Sources, New York State Department of Health, Division of Environmental Protection, Bureau of Public Water Supply Protection, 1982.
- 16. Telecon Karen A. Hartnett, URS Corporation, to Fire Chief, City of Utica, September 2, 1987.
- 17. 1980 Census of Population, Number of Inhabitants, New York, United States Department of Commerce, Bureau of the Census.

HARBOR POINT PROPERTY LAND INVESTIGATIONS STEP 3 LAND REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York

HARBOR POINT PROPERTY RIVER INVESTIGATIONS STEP 4 RIVER REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York

REF. 3

Uncontrolled Hazardous Waste Site Ranking System

A Users Manual (HW-10)

Originally Published in the July 16, 1982, Federal Register

United States Environmental Protection Agency

TABLE 2
PERMEABILITY OF GEOLOGIC MATERIALS*

TYPE OF MATERIAL	APPROXIMATE RANGE OF HYDRAULIC CONDUCTIVITY	ASSIGNED VALUE
Clay, compact till, shale; unfractured metamorphic and igneous rocks	< 10 ⁻⁷ cm/sec	0
Silt, loess, silty clays, silty loams, clay loams; less permeable limestone, dolomites, and sandstone; moderately permeable till	<10 ⁻⁵ \geq 10 ⁻⁷ cm/sec	1
Fine sand and silty sand; sandy loams; loamy sands; moderately permeable limestone, dolomites, and sandstone (no karst); moderately fractured igneous and metamorphic rocks, some coarse till	<10 ⁻³ ≥ 10 ⁻⁵ cm/sec	2
Gravel, sand; highly fractured igneous and metamorphic rocks; permeable basalt and lavas; karst limestone and dolomite	>10 ⁻³ cm/sec	3

^{*}Derived from:

Davis, S. N., Porosity and Permeability of Natural Materials in Flow-Through Porous Media, R.J.M. DeWest ed., Academic Press, New York, 1969

Freeze, R.A. and J.A. Cherry, Groundwater, Prentice-Hall, Inc., New York, 1979

TABLE 3

CONTAINMENT VALUES FOR GROUND WATER ROUTE

Assign containment a value of 0 if: (1) all the hazardous substances at the facility are underlain by an essentially non permeable surface (natural or artificial) and adequate leachate collection systems and diversion systems are present; or (2) there is no ground water in the vicinity. The value "0" does not indicate no risk. Rather, it indicates a significantly lower relative risk when compared with more serious sites on a national level. Otherwise, evaluate the containment for each of the different means of storage or disposal at the facility using the following guidance.

A. Surface Impoundment		C. Pilen	
	Assigned Value		salgued Value
Sound run-on diversion structure, exactificial) compatible with the waste, and	0	Piles uncovered and waste stabilized; or piles covered, waste unstabilized, and essentially non permeable liner	G
adequate leachate collection system Ensentially non permeable compatible liner with no leachate collection system; or	ı	Piles uncovered, waste unstablized, moderately permeable liner, and leachate collection system	L
inadequate freeboard Potentially unnound run-on diversion	2	Piles uncovered, waste unstabilized, moderately permeable liner, and no leachate collection system	2
atructure; or moderately permeable compatible liner		Piles uncovered, waste unstablized, and no liner	3
Unsound run-on diversion atructure; no liner; or incompatible liner	3	D. Landfill	
B. Containers		<u>.</u>	Assigned Value
	Assigned Value	Essentially non permeable liner, liner	0
Containers scaled and in sound condition, adequate liner, and adequate leschate	0	compatible with waste, and adequate leachate collection system	
collection system Containers sealed and in sound condition, no liner or moderately permeable liner	1	Essentially non permeable compatible liner, no leachate collection system, and landfill surface precludes ponding	1
Containers leaking, moderately permeable	2	Moderately permeable, compatible liner, and landfil surface procludes pending	P1 2
Containers leaking and no liner or incompatible liner	3	No liner or incompatible liner; moderately permeable compatible liner; landfill nurface encourages ponding; no run-on control	3

TABLE 4 WASTE CHARACTERISTICS VALUES FOR SOME COMMON CHEMICALS

ander/agean				
Acotaldulydo		•	•	2
Acetic Acid] 3]	•	2	1
Acotons	2	0	3	•
Aldrin	3	3	1	•
Americ, Askydrous	3	•	1	•
<u>Antilip</u>	3	1	*	•
Bussens	3	1	3	•
Carbon Totroshloride	1	3	•	•
Chleriene		•	•	•
Chieroteneone		2	3	•
Caleroform	3	3	•	•
Greenl-0	3	1	3	•
Crossl-WP	3	1	1	•
Cyclohomes	1	2	3	•
Bairta	. 3		1	•
Ruly 1 Russia			3	
Portsaldskyde	,		3	
Poznác Asiá Brisvehlesie Asiá			•	
Loopropyl Ether	1	1	3	
Linimo	15	,	i	
Matheire		1	3	
Notical Exteri Second			,	
Methyl Perethies in Sylene Selection		4	- 1	2
Bubthelese		1		
Mitrie Asid				
Incodes .		٠,		2
300		,		
	,	1	•	•
Petralous, Escarina (Pust OLL Sp. 1)	'	١.	•	•
Property.	2	1	2	•
Pulfurio Acid	3	•	•	2
Teluçus	2	1	3	•
Trichistorium	1 2	3	3	•
et-Trichlercethine	2	2	1	•
Zylens		1	3	•

Tex. W. I., Innervee Presertion of Industrial Interials, Van Nestrand Shaisheld Co., New York, 4th ed., 1975. The highest rating lioted under each chanical is used.

²JRS Association, Inc., <u>Hethedelows for Retine the Suseri</u> <u>Princial of these Dismont Situs</u>, May 3, 1960.

Stational Pire Protestion Accosistion, National Pire Godos, Vol. 13, No. 49, 1977.

^{*}Professional judgment based on information contained in the U.S. Cocot Guard CRIS linguisting Chesical Sate, 1978.

A Professional judgment based on egisting literature.

TABLE 5

PERSISTENCE (BIODEGRADABILITY) OF SOME ORGANIC COMPOUNDS*

VALUE - 3	STORTA SENSTREES CONCORDS
aldrin benzopyrene benzothizole benzothizole benzothizolene benzothizolene benzothizolene benzothizolene brouschierebenzene brouschierebenzene chloredere betzenl brouschierelene chloredere benzeghrinene bis-chlereinspezighri ether n-chlorenterebenzene non non diburyl phthalate 1, 4-dichlerebenzene dichlorediflusvesthene dichtrin phthalate 1, delderen	heptachler heptachler openide 1,2,3,3,7,7-heptachlerenethernene henochlerebensene henochlerebensene henochleresel,3-betachlene henochlereselenene henochleresthene methyl henochlesele pentachleresphenel 1,1,3,3-testrochlerescetene tetrachleresphenel titmattylbeneothlesele trichleresphenel trichleresphenel trichleresphenel trichleresphenel trichleresphenel trichleresphenel triphonyl trichleresphenel triphonyl phesphene
dictryl pithelete dife-ethylhenyl)phthelete dihenyl pithelete di-losbutyl pithelete dimethyl pithelete d,-ddinitro-3-milesphesel dipropyl pithelete endrin	boundern esthen tettenhieride chierofern chierofern chierontichieronthene dibroundichieronthene totrechieronthene 1,1,2-trichieronthene

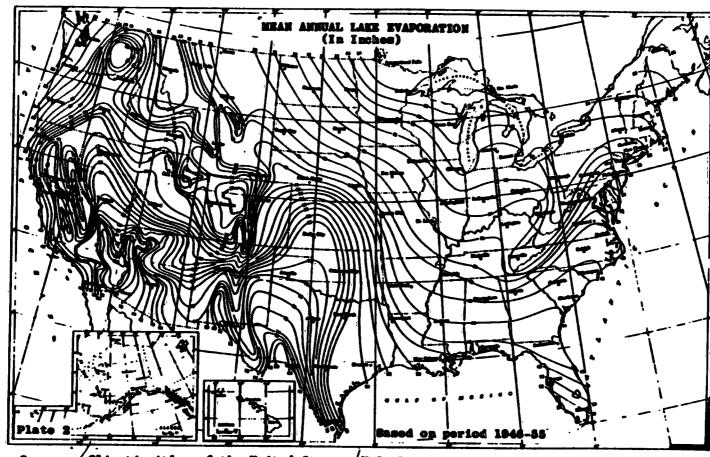
acatyless dichloride behenic acid, methyl ester beaness beaness beaness oulfenie ecid butyl beaness butyl brounde e-caprolectes ester-dizelfide e-caprolectes (access 1,2-dichlorosthess 1,2-dinethyl meghtheless 1,3-dimethyl meghtheless 1,4-dimethyl meghtheless 1,4-dimethyl edipets p-dedocase ethyl beaness 2-ethyl-p-beanes 2-ethyl-p-beanes	lineases sethyl ester of lignoseric or methon 2-methyl-5-othyl-pyridine - methyl nephthelene methyl planitate methyl planitate methyl carbinel methyl carbinel methyl descrete mphthelene mesene methologic methologic methologic methologic methologic methologic methologic methologic methologic perpylleneme l-explanel
o-ethyitelesses	telecie
leedeesse	vizzi beneze
leepruphyi bessesse	Aylene

	MTHE - 3	PRESENT CONTINUE
acceaphthylene etrenine (diethyl) etrenine berbital berneel bromelene campter chlorebeneene 1,2-bis-chloresthys et b-chloresthyl enter chloresthyl ether chloresthyl ether di-t-butyl-p-beneesian dichloresthyl ether dit-t-butyl-p-beneesian dichloresthyl ether dihyrocarvene dinethyl eulfenide 2,6-dinitrotolusse	her T	cio-l-cthyl-4-asthyl-1, 3-dienolano tran-2-cthyl-4-asthyl-1, 3-dienolano gusiacol 3-bytemyodipositrile isopheroso indene isoberosol isopropayl-r-isopropyl bessene 2-asthony bighanyl enthyl bighanyl enthyl bighanyl enthyl bighanyl enthyl indene mothylindene mothylindene mitroanisola ultroanisola ultroanisola trinothyl-triono-bennhydro-trionina trinothyl-triono-bennhydro-trionina iomar

TALES - 8	SOMPERSTATION CONFIGURE
acestaldehyda acestic acid acestic acid acestica acestican acestic	enthyl bennoste 3-enthyl betseel enthyl ethyl heteen 2-enthylpropenel ectadecane pentadecane pentadecane pentadecane pentadecane pentadecane perspientane tetradecane o-tridecane n-undecane

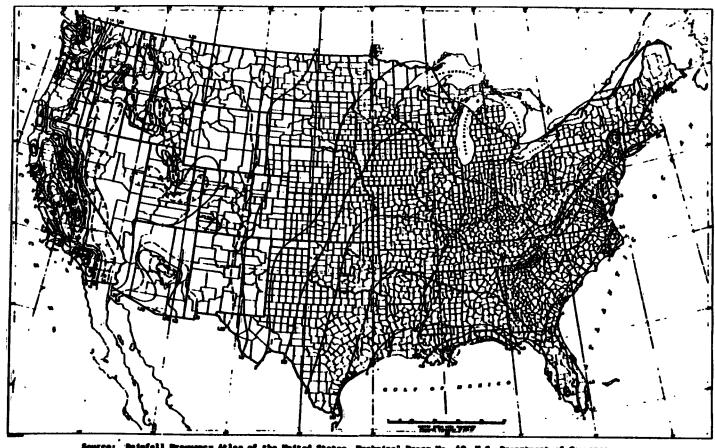
Assign containment a value of 0 if: (1) all the waste at the site is surrounded by diversion structures that are in sound condition and adequate to contain all runoff, spills, or leaks from the waste; or (2) intervening terrain precludes runoff from entering surface water. Otherwise, evaluate the containment for each of the different means of storage or disposal at the site and assign a value as follows:

A. Surface Impoundment		C. Mapte Piles.	
<u>Aus1</u>	gned Value		Ventaucd Anim
Sound diving or diversion structure, adequate freeboard, and no erosion evident	0	Piles are covered and surrounded by sound diversion or containment system	0
Sound diking or diversion structure, but insdequate freeboard	1	Piles covered, vastes unconsolidated, diversion of containment system not adequate	1
Diking out leaking, out potentially unsound	2	Piles not covered, wastes unconsoli- dated, and diversion or containment system potentially unsound	2
Diking unsound, leaking, or in danger of cullapse	3	Piles not covered, wastes unconsolidated, and no diversion or containment or diversion	j
B. Comtainara		mystem leaking or in danger or collapse	
Asot	ned Value	P. Landfill	
Containers sealed, in sound condition, and our- rounded by sound diversion or containment system	0	·	Applymed Vator
Containers sealed and in sound condition, but not surrounded by sound diversion or containment system	4	Landfill slope precludes runoff, landfill surrounded by sound diversion system, or landfill has adequate cover material	0
Containers leaking and diversion or containment structures potentially unsound	2	Laudfill not adequately covered and diversion system sound	1
Containers leaking, and up diversion or containment structures or diversion structures leaking or in	3	Landfill not covered and diversion system potentially unsound	2
dunger of collapse		Landfill not covered and no diversion system present, or diversion system unnound	3



Source: Climatic Atlas of the United States, U.S. Department of Commerce, Hational Climatic Cemter, Ashville, N.C., 1979.

FIGURE 4
MEAN ANNUAL LAKE EVAPORATION
(IN INCHES)



Source: Reinfell Progressy Atlas of the United States, Technical Paper No. 40, U.S. Department of Councres, U.S. Government Printing Office, Unchington, D.C., 1963.

FIGURE 8 1-YEAR 24-HOUR RAINFALL (INCHES)

Ground Water Use (continued)	Assigned Value
Drinking water with municipal water from alternate unthreatened sources presently available (i.e., minimal hookup requirements); or commercial, industrial or irrigation with no other water source presently available	2
Drinking water; no municipal water from alternate unthreatened sources presently available	3

Distance to nearest well and population served have been combined in the matrix below to better reflect the important relationship between the distance of a population from hazardous substances and the size of the population served by ground water that might be contaminated by those substances. To determine the overall value for this combined factor, score each individually as discussed below. Match the individual values assigned with the values in the matrix for the total score.

Value for Population		Value fo	r Distance t	o Nearest	Well
Served	. 0	1	2	3	4
0	0	0	0	Ø	0
1	0	4	6	8	10
2	0	8	12	16	20
3	0	12	18	24	30
4	0	16	24	32	35
5	0	20	30	35 `	40

Distance to nearest well is measured from the hazardous substance (not the facility boundary) to the nearest well that draws water from the aquifer of concern. If the actual distance to the nearest well is unknown, use the distance between the hazardous

substance and the nearest occupied building not served by a public water supply (e.g., a farmhouse). If a discontinuity in the aquifer occurs between the hazardous substance and all wells, give this factor a score of 0, except where it can be shown that the contaminant is likely to migrate beyond the discontinuity. Figure 6 illustrates how the distance should be measured. Assign a value using the following guidance:

Population served by ground water is an indicator of the population at risk, which includes residents as well as others who would regularly use the water such as workers in factories or offices and students. Include employees in restaurants, motels, or campgrounds but exclude customers and travelers passing through the area in autos, buses, or trains. If aerial photography is used, and residents are known to use ground water, assume each dwelling unit has 3.8 residents. Where ground water is used for irrigation, convert to population by assuming 1.5 persons per acre of irrigated land. The well or wells of concern must be within three miles of the hazardous substances, including the area of known aquifer contamination, but the "population served" need not be. Likewise, people within three miles who do not use water from the aquifer of concern are not to be counted. Assign a value as follows:

(3)

<u>Population</u>	Assigned	Value
0	0	
1-100	1	
101-1,000	2	
1,001-3,000	3	
3,001-10,000	4	
> 10,000	5	

Dangerous Properties of Industrial Materials

Sixth Edition

N. IRVING SAX

Assisted by:

Benjamin Feiner/Joseph J. Fitzgerald/Thomas J. Haley/Elizabeth K. Weisburger

pisaster Hazard: Dangerous; shock will explode it; when heated, burns and emits acrid fumes; can react on contact with oxidizing materials.

NAPHTHA, COAL TAR

CAS RN: 8030306

NIOSH #: OI 9450000

Dark straw-colored to colorless liquid. Sol in benzene, toluene, xylene, etc. bp: 149°-216°, flash p: 107°F (CC), d: 0.862-0.892, autoign. temp.: 531°F.

SYNS:

BI NZIN 160 DEGREE BENZOL NAPHTHA NAPHTHA, PETROLEUM PETROLEUM BENZIN PETROLEUM NAPHTHA

COAL TAR NAPHTHA DISTILLATE I IGHT LIGROIN

NAITA (POLISH)

TOXICITY DATA: ini-rat LCLo: 1600 ppm/6H CODEN: CHINAG 17,1078,39

TLV: Air: 300 ppm DTLVS* 4,433,80. OSHA Standard: Air: TWA 100 ppm (SCP-G) FEREAC 39,23540,74. "NIOSH Manual of Analytical Methods" VOL 2 S86. Reported in EPA TSCA Inventory, 1980.

THR: MOD via inhal route. Can cause unconsciousness which may go to coma, stentorious breathing and bluish tint to the skin. Recovery follows removal from exposure. In mild form, intoxication resembles drunkenness. On a chronic basis no true poisoning; sometimes headache, lack of appetite, dizziness, sleeplessness, indigestion and nausea. A common air contaminant. See oils.

Fire Hazard: Mod, when exposed to heat or flame; can react with oxidizing materials. Keep containers tightly closed.

Explosion Hazard: Slight.

To Fight Fire: Foam, CO2, dry chemical.

alpha-NAPHTHAL

CAS RN: 66773

NIOSH #: QJ 0175000

TOXICITY DATA:

CODEN:

wu-dog LDLo:330 mg/kg ZMWIAJ 19,545,1881

Reported in EPA TSCA Inventory, 1980. TUR: HIGH scu.

NAPHTHALENE

CAS RN: 91203

NIOSH #: QJ 0525000

mf: C10H8; mw: 128.18

Aromatic odor, white, crystalline, volatile flakes. mp: 80.1°, bp: 217.9°, flash p: 174°F (OC), d: 1.162, lel = 0.9%, uel = 5.9%, vap. press: 1 mm @ 52.6°, vap. d: 4.42. Autoign temp: 1053°F (567°C); sol in alc, benzene. Insol in water; very sol in ether, CCl4, CS2 hydronaphthaknes, in fixed and volatile oils.

SYNS:

(AMPHOR TAR WITH BALLS WITH ILAKES MAPTALEN (POLISH)

*APHTHALINE

NAPHTHENE NCI-C52904 TAR CAMPHOR WHITE TAR

TOXICITY DATA: ipr-rat TDLo:5925 mg/kg (1-15D

skn-rbt 495 mg open MLD eye-rbt 100 mg MLD

scu-rat TDLo:3500 mg/kg/12W-

I:ETA orl-chd LDLo: 100 mg/kg unk-man LDLo:74 mg/kg orl-rat LD50:1780 mg/kg ipr-mus LD50: 150 mg/kg scu-mus LD50:969 mg/kg

ivn-mus LD50:100 mg/kg orl-dog LDLo: 400 mg/kg orl-cat LDLo: 1000 mg/kg orl-rbt LDLo:3 gm/kg orl-mam LD50:1000 mg/kg CODEN:

TXAPA9 48,A35,79

UCDS** 1/11/68 BIOFX* 16-4/70 APAVAY 329,141,56

28ZRAQ -,228,60 85DCAI 2.73.70 BIOFX* 16-4/70 NTIS** AD691-490 TOIZAG 20(5/6),772,73 CSLNX* NX#00203 HBAMAK 4,1289,35 HBAMAK 4,1289,35 HBAMAK 4,1289,35 FMCHA2 -, D213,80

Aquatic Toxicity Rating: TLm96:10-1 ppm WQCHM* 3,-,74. TLV: Air: 10 ppm DTLVS* 4,293,80. Toxicology Review: 38ZNAA 1(1),93,71; JOPDAB 59,1,61; 27ZTAP 3,30,69. OSHA Standard: Air: TWA 10 ppm (SCP-T) FEREAC 39,23540,74. DOT-ORM-A, Label: None FEREAC 41,57018,76. Currently Tested by NTR for Carcinogenesis by Standard Bioassay Protocol as of Sept 1980. "NIOSH Manual of Analytical Methods" VOL 3 S292. Reported in EPA TSCA Inventory,

THR: MOD orl and HIGH ipr, ivn. An exper ETA. May be used as an insecticide. Systemic reactions include nausea, headache, diaphoresis, hematuria, fever, anemia, liver damage, vomiting, convulsions and coma. Poisoning may occur by ing of large doses, inhal or skn absorption.

Fire Hazard: Mod, when exposed to heat or flame; reacts with oxidizing materials. Reacts violently with CrO₃.

Spontaneous Heating: No.

Explosion Hazard: Mod, in the form of dust, when exposed to heat or flame.

To Fight Fire: Water, CO2, dry chemical.

Incomp: Dinitrogen pentaoxide.

1-NAPHTHALENEACETAMIDE

CAS RN: 86862

NIOSH #: QJ 0590000

mf: C₁₂H₁₁NO; mw: 185.24

SYNS:

NAPHTHALENE ACETAMIDE ALPHA-NAPHTHALENEACET-

ALPHA-NAPHTHYLACETAMIDE 1-NAPHTHYLACETAMIDE

AMIDE

TOXICITY DATA: orl-mam LD50:1000 mg/kg 2 CODEN: FMCHA2 -, D143,75

Reported in EPA TSCA Inventory, 1980.

THR: MOD orl.

Disaster Hazard: When heated to decomp it emits tox fumes of NO_x.

1-NAPTHTALENEACETIC ACID

CAS RN: 86873

NIOSH #: QJ 0875000

 $mf: C_{12}H_{10}O_2; mw: 186.22$



1882 2-METHYLNAPHTHALENE

TOXICITY DATA:

1 CODEN:

mma-sat 6 mmol/L/2H orl-rat LDLo:5000 mg/kg CNREA8 39,4152,79 28ZRAQ -,55,60

Reported in EPA TSCA Inventory, 1980. EPA TSCA 8(a) Preliminary Assessment Information Proposed Rule FERREAC 45,13646,80.

THR: MUT data. LOW orl. See also 2-methyl naphtha-

Fire Hazard: Mod, via heat, flames, oxidizers.

To Fight Fire: Dry chemical, CO₂, water spray or mist, foam.

Disaster Hazard: When heated to decomp it emits acrid smoke and fumes.

2-MÉTHYLNAPHTHALENÉ

CAS RN: 91576

NIOSH #: QJ 9635000

mf: C₁₁H₁₀; mw: 142.21

Solid, insol in water, sol in alc and ether, d: 1.0058 @ 20°/4°, bp: 241.1°, mp: 34.58°.

SYN: BETA-METHYLNAPHTHALENE

TOXICITY DATA:

CODEN:

orl-rat LDLo:5000 mg/kg

28ZRAQ -,55,60

Reported in EPA TSCA Inventory, 1980. EPA TSCA 8(a) Preliminary Assessment Information Proposed Rule FERREAC 45,13646,80

THR: LOW orl.

Disaster Hazard: When heated to decomp it emits acrid smoke and fumes.

2-METHYL-1,4-NAPHTHALENEDIOL

CAS RN: 481856

NIOSH #: QJ 5075000

mf: C₁₁H₁₀O₂; mw: 174.21

SYNS:

METHYLNAPHTHOHYDRO-

2-METHYL-1,4-NAPHTHOQUINOL

QUINONE

2-METHYL-1,4-NAPHTHOHYDRO-

QUINONE

TOXICITY DATA:

3

CODEN: JPETAB 75,111,42

orl-mus LD50:30 mg/kg ipr-mus LDLo:400 mg/kg

CRSBAW 143,585,49

THR: HIGH orl, ipr.

Disaster Hazard: When heated to decomp it emits acrid

smoke and fumes.

2-METHYL-1,4-NAPHTHALENEDIOL DIBUTYRATE

CAS RN: 53370448

NIOSH #: OJ 5255000

mf: $C_{19}H_{22}O_4$; mw: 314.41

SYNS:

2-METHYL-1,4-NAPHTHOHYDRO-CHINON-DIBUTYRAT (GER-

VITAMIN K4-DIBUTYRAT (GER-

MAN)

MAN)

TOXICITY DATA:

2-1 CODEN:

ori-mus LD50:12 gm/kg ipr-mus LD50:1 gm/kg

ARZNAD 17.1339.67 ARZNAD 17,1339,67 THR: MOD ipr; LOW orl.

Disaster Hazard: When heated to decomp it emits acrid

smoke and fumes.

P-METHYL-N-(1-NAPHTHALENYLOXY) PHOSPHONAMIDOTHIOIC ACID-O-ETHYL **ESTER**

CAS RN: 63815532

NIOSH #: SZ 6260000

mf: C₁₃H₁₆NO₂PS; mw: 281.33

SYN: BAY 39197

TOXICITY DATA:

CODEN:

orl-rat LD50:250 mg/kg orl-bwd LD50:18 mg/kg

TXAPA9 21,315,72 TXAPA9 21,315,72

THR: HIGH orl.

Disaster Hazard: When heated to decomp it emits very

tox fumes of SO_x , PO_x and NO_x .

5-METHYLNAPHTHO(1,2,3,4-def)CHRYSENE

CAS RN: 2869092

NIOSH #: QL 0350000

mf: C₂₅H₁₆; mw: 316.41

SYN: 2'-METHYL-1,2:4,5-DIBENZOPYRENE

TOXICITY DATA:

CODEN:

scu-mus TDLo:72 mg/kg/9W-I:ETA COREAF 259,3899,64

THR: An exper ETA.

Disaster Hazard: When heated to decomp it emits acrid

smoke and fumes.

6-METHYLNAPHTHO(1,2,3,4-def)CHRYSENE

CAS RN: 2869105

NIOSH #: QL 0525000

mf: C₂₅H₁₆; mw: 316.41

SYN: 3'-METHYL-1.2:4.5-DIBENZOPYRENE

TOXICITY DATA:

CODEN:

scu-mus TDLo:72 mg/kg/9W-I:ETA

COREAF 259,3899,64

THR: An exper ETA.

Disaster Hazard: When heated to decomp it emits acrid

smoke and fumes.

2-METHYL-1,4-NAPHTHOQUINONE

CAS RN: 58275

NIOSH #: QL 9100000

mf: C₁₁H₈O₂; mw: 172.19

SYNS:

KIPCA, OIL SOLUBLE 2-METHYL-1,4-NAPHTHOCHINON

THYLOQUINONE **USAF EK-5185** VITAMIN K3

(GERMAN)

3-METHYL-1,4-NAPHTHOQUI-

NONE

CODEN: **TOXICITY DATA:** 3 ABCHA 6,45,327,81

mmo-sat 4 ug/plate mmo-omi-6 ppm mma-sat 30 nmol/plate

skn-mus TDLo: 1860 mg/kg/27W-I:ÉTA

skn-mus TD:8400 mg/kg/21W-I:ETA

ipr-rat LD50:75 mg/kg ivn-rat LD50:800 mg/kg BBRCA9 94,737,80 BJCAAI 7,482,53 CNREA8 19,413,59

POASAD 34,114,53

TXAPA9 18,185,71 ARZNAD 18,678,68

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BENZEDRINE SULFATE

CAS RN: 156310 NIOSH #: SI 1225000

mf: $C_{18}H_{26}N_2 \cdot H_2O_4S$; mw: 368.54

SYNS:

PHENETHYLAMINE, ALPHA-DL-ALPHA-METHYLPHENETH-METHYL-, SULFATE (2:1) YLAMINE SULFATE DIAMPHETAMINE SULFATE 1-PHENYL-2-AMINOPROPANE SULFATE

TOXICITY DATA: ipr-rat LDLo:25 mg/kg

CODEN: JPETAB 100,267,50 scu-rat LDLo: 10 mg/kg JPETAB 71,62,41 scu-mus LD50:14 mg/kg JPETAB 87,214,46 ipr-gpg LDLo: 50 mg/kg JPETAB 100,267,50

THR: HIGH ipr, scu. See also sulfates. Disaster Hazard: When heated to decomp it emits very tox fumes of SO_x and NO_x .

D-BENZEDRINE SULFATE

CAS RN: 51638 NIOSH #: SI 1400000 mf: C₁₈H₂₆N₂•H₂O₄S; mw: 368.54

SYNS:

AMPHEDRINE **DEXTROAMPHETAMINE SULFATE AMPHEREX** DEXTRO-ALPHA-METHYLPHENE-(+)-AMPHETAMINE SULFATE THYLAMINE SULFATE D-AMPHETAMINE SULFATE ORANGES DEXAMPHETAMINE SULFATE PHENEDRINE DEXAMYL PHENOPROMIN DEXEDRINA D-1-PHENYL-2-AMINOPROPANE DEXEDRINE SULFATE SULFATE DEXTRO-1-PHENYL-2-AMINO-D-ALPHA-METHYLPHENETHYL-PROPANE SULFATE AMINE SULFATE D-BETA-PHENYLISOPROPYL-OBESEDRIN AMINE SULFATE **FASTBALLS** DEXTRO-BETA-PHENYLISOPRO-HEARTS **PYLAMINE SULFATE**

TOXICITY DATA: ipr-mus TDLo: 50 mg/kg/(8D preg):TER unk-mus TDLo:50 mg/kg/(8D preg):TER orl-rat LD50:38 mg/kg ipr-rat LD50:70 mg/kg scu-rat LD50:200 mg/kg ivn-rat LD50:30 mg/kg orl-mus LD50:33 mg/kg ipr-mus LD50:72 mg/kg scu-mus LD50:16 mg/kg ivn-mus LD50:30 mg/kg

orl-dog LD50:10 mg/kg

ivn-dog LD50:3 mg/kg

ivn-rbt LD50:10 mg/kg

TJADAB 1.413.68 TJADAB 1.413.68 JOPDAB 69,663,66 TXAPA9 45(1),49,78 12VXA5 8,335,68 JPETAB 110,180,54 TXAPA9 21,302,72 JPETAB 128,176,60 AIPTAK 184,34,70 JPETAB 137,365,62 PSEBAA 118,557,65 PSEBAA 118,557,65

JPETAB 110,180,54

CODEN:

Toxicology Review: ISYAM* -,343,70; 27ZTAP 3,46,69. THR: An exper TER. HIGH orl, ipr, scu, ivn. A habitforming stimulant. See also sulfates.

Disaster Hazard: When heated to decomp it emits very tox fumes of SO_x and $NO_{\hat{x}}$.

L-BENZEDRINE SULFATE

CAS RN: 51627 NIOSH #: SI 1575000 mf: C₁₈H₂₆N₂•H₂O₄S; mw: 368.54

(-)-AMPHETAMINE SULFATE L-1-PHENYL-2-AMINOPROPANE L-AMPHETAMINE SULFATE SULFATE LEVEDRINE

TOXICITY DATA: 3 CODEN: scu-rat LDLo: 160 mg/kg JPETAB 71,62,41 ipr-mus LD50:232 mg/kg JPETAB 158,135,67

THR: HIGH scu, ipr. See also sulfates. Disaster Hazard: When heated to decomp it emits very tox fumes of SO_x and NO_x .

BENZENAMINE HYDROCHLORIDE

CAS RN: 142041 NIOSH #: CY 0875000 mf: C₆H₇N•ClH; mw: 129.60

Crystals. vap. d: 4.46, d: 1.22, mp: 198°, bp: 245°, flash p: 380°F (OC).

SYNS:

ANILINE HYDROCHLORIDE CHLORID ANILINU (CZECH) "ANILINE SALT" NCI-CO3736 CHLORHYDRATE D'ANILINE 'USAF EK-442 (FRENCH)

TOXICITY DATA: 3 CODEN: skn-rbt 500 mg/24H MOD 28ZPAK -,65,72 eye-rbt 20 mg/24H SEV 28ZPAK -,65,72 orl-rat TDLo: 130 gm/kg/2Y-NCITR* NCI-CG-TR-C:CARC 130,78 orl-rat TD:238 gm/kg/2Y-C:CARC NCITR* NCI-CG-TR-130.78 orl-rat LD50:1072 mg/kg NTIS** PB214-270 ipr-rat LDLo:500 mg/kg NCNSA6 5,11,53 orl-mus LD50:841 mg/kg NTIS** PB214-270 ipr-mus LD50:300 mg/kg NTIS** AD277-689 orl-rat TD:137 gm/kg/60W-C:ETA IARC** 27,39,82 orl-rat TD:2163 gm/kg/2Y-C:CAR IARC** 27,39,82 orl-rat TD:4326 gm/kg/2Y-C:CAR IARC** 27,39,82

Aquatic Toxicity Rating: TLm96: 100-10 ppm WQCHM* 2,-,74. NCI Carcinogenesis Bioassay Completed; Results Positive: Rat (NCITR* NCI-CG-TR-130,78). NCI Carcinogenesis Bioassay Completed; Results Negative: Mouse (NCITR* NCI-CG-TR-130,78). Reported in EPA TSCA Inventory, 1980. EPA TSCA 8(a) Preliminary Assessment Information Proposed Rule FERREAC 45,13646,80.

THR: An exper CARC, HIGH ipr; MOD orl, ipr. MOD skn irr, SEV eye irr in rbt. See also aniline.

Fire Hazard: Slight, when exposed to heat or flame. Spontaneous Heating: No.

Disaster Hazard: Dangerous; when heated to decomp, or on contact with acid or acid fumes, emits highly toxic fumes of aniline and chlorine compounds; can react vigorously with oxidizing materials.

To Fight Fire: Water, CO2, water mist or spray, dry chemical.

BENZENE

CAS RN: 71432 NIOSH #: CY 1400000 mf: C₆H₆; mw: 78.12

Clear colorless liquid. mp: 5.51°, bp: 80.093°-80.094°, flash p: 12°F (CC), d: 0.8794 @ 20°, autoign. temp.: 1044°F, lel: 1.4%, uel: 8.0%, vap. press: 100 mm @ 26.1°, vap. d: 2.77, ulc: 95-100.

SYNS:
(6)ANNULENE
BENZEEN (DUTCH)
BENZEN (POLISH)
BENZOL
BENZOLENE
BENZOLO (ITALIAN)
BICARBURET OF HYDROGEN
CARBON OIL

COAL NAPHTHA
CYCLOHEXATRIENE
FENZEN (CZECH)
MINERAL NAPHTHA
MOTOR BENZOL
NCI-C55276
PHENYL HYDRIDE
PYROBENZOLE

TOXICITY DATA: 3
skn-rbt 15 mg/24H open MLD
eye-rbt 88 mg MOD
eye-rbt 2 mg/24H SEV
cyt-rat-scu 12 gm/kg/12D-I
mnt-mus-ipr 500 uL/kg
cyt-mus-orl 100 uL/kg
cyt-mus-ipr 100 uL/kg
dlt-mus-ipr 5 mg/kg
cyt-rbt-scu 8400 mg/kg
scu-mus TDLo:2700 mg/kg/(13D
preg): TER

scu-mus TDLo:2700 mg/kg/(13D preg):TER
ihl-hmn TCLo:100 ppm/10Y-I:CAR
orl-rat TDLo:52 gm/kg/52W-I:CAR
skn-mus TDLo:1200 gm/kg/
49W-I:NEO
scu-mus TDLo:600 mg/kg/

17W-I:ETA
par-mus TDLo:670 mg/kg/
19W-I:ETA

19W-I:ETA
ihl-hmn TC:400 ppm/8Y-I:ETA
thl-man TC:2100 mg/m3/4Y-I:CAR
orl-rat TD:10 gm/kg/52W-I:CAR
orl-hmn TDLo:130 mg/kg:CNS
thl-hmn LCLo:20000 ppm/5M
thl-hmn TCLo:210 ppm:BLD
ihl-rat TCLo:670 mg/m3/24H (15D
pre/1-22D preg)
thl-rat TCLo:56600 ug/m3/24H
(1-22D preg)

thl-rat TCLO:50 ppm/24H (7-14D preg)
ihl-rat TCLO:150 ppm/24H (7-14D preg)

scu-mus TDLo:1100 mg/kg (12D preg) scu-mus TDLo:2700 mg/kg/(13D

preg) TFX:TER
orl-mus TDLo:9 gm/kg (6-15D preg)
orl-mus TDLo:12 gm/kg (6-15D preg)
orl-rat TD:10 gm/kg/52W-I

TFX:CAR thl-hmn TCLo: 100 ppm: CNS unk-man LDLo: 194 mg/kg orl-rat LD50:3800 mg/kg ihl-rat LC50:10000 ppm/7H ipr-rat LDLo: 1150 mg/kg orl-mus LD50:4700 mg/kg ihl-mus LC50:9980 ppm ipr-mus LD50:990 ug/kg ori-dog LDLo: 2000 mg/kg thi-dog LCLo: 146000 mg/m3 ihl-cat LCLo: 170000 mg/m3 ivn-rbt LDLo:88 mg/kg ipr-gpg LDLo: 527 mg/kg scu-frg LDLo: 1400 mg/kg ihl-mam LCLo:20000 ppm/5M CODEN:
AIHAAP 23,95,62
AMIHAB 14,387,56
28ZPAK -,23,72
GTPZAB 17(3),24,73
ENMUDM 2,43,80
ENMUDM 2,43,80
ENMUDM 2,43,80
ENMUDM 2,43,80
TPKVAL 15,30,79
PSDTAP 15,275,74
AMBNAS 17,285,70

TRBMAV 37,153,78 MELAAD 70,352,79 BJCAAI 16,275,62

KRANAW 9,403,32

KLWOAZ 12,109,33

BLOOAW 52,285,78 NEJMAG 271,872,64 MELAAD 70,352,79 AHYGAJ 31,336,1897 29ZUA8 -,-,53 27ZXA3 -,341,63 HYSAAV 33,327,68

HYSAAV 33,112,68

JHEMA2 24,363,80

JHEMA2 24,363,80

TOXID9 1,125,81

AMBNAS 17,285,70

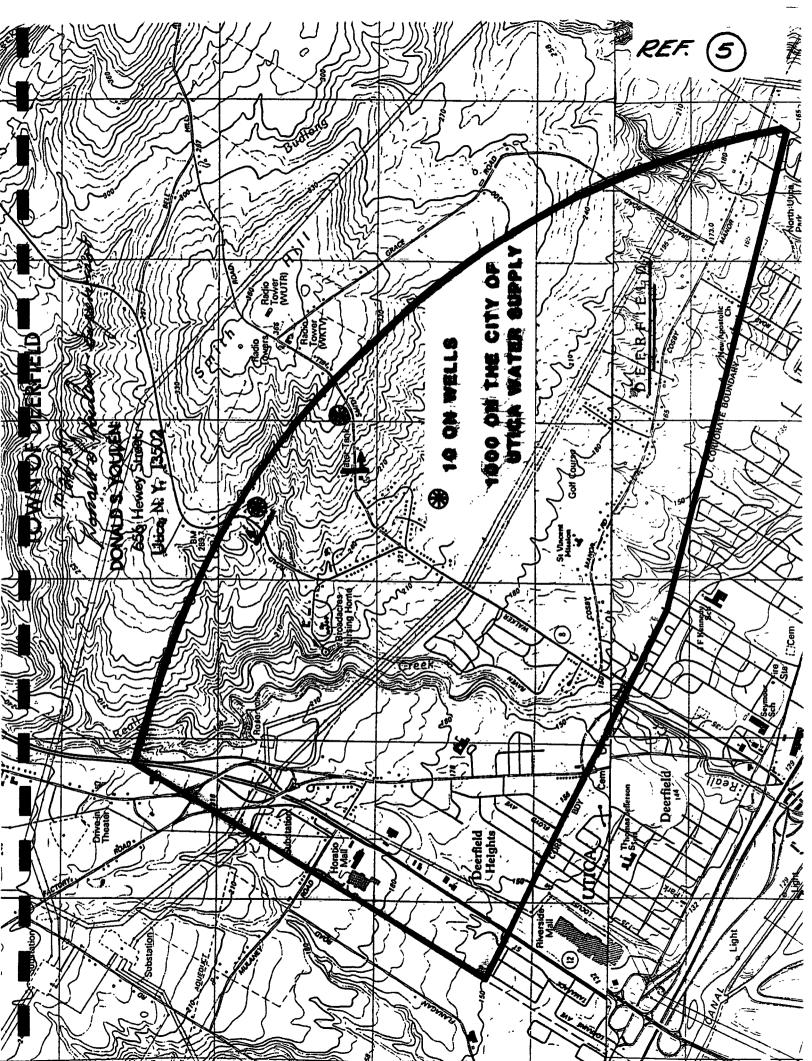
TJADAB 19,41A,79 TJADAB 19,41A,79 MELAAD 70,352,79

INMEAF 17,199,48
85DCAI 2,73,70
TXAPA9 19,699,71
28ZRAQ -,113,60
TXAPA9 1,156,59
HYSAAV 32,349,67
JIHTAB 25,366,43
AGGHAR 18,109,60
HBAMAK 4,1313,35
HBTXAC 1,324,56
HBTXAC 1,324,56
JTEHD6 -(Suppl.2),45,77
HBTXAC 1,42,56
HBAMAK 4,1313,35
AEPPAE 138,65,28

Aquatic Toxicity Rating: TLm96: 100-10 ppm WQCHM* 2,-,74. Carcinogenic Determination: Human Suspected IARC** 7,203,74.

TLV: Air: 10 ppm DTLVS* 4,37,80. Toxicology Review: ARPAAQ 11,434,31; EVHPAZ 11,163,75; AEHLAU 22,373,71; PAREAQ 4,1,52; FNSCA6 2,67,73; MU-REAV 47(2),75,78; AMSVAZ 118,354,44; ZHPMAT 166,113,78; JTEHD6 -(suppl.2),69,77; PHRPA6 41,1357,26; CTOXAO 11,531,77; BNYMAM 54, 413,78; KRANAW 9,403,32; 27ZTAP 3,22,69. OSHA Standard: Air: TWA 10 ppm; CL 25 ppm; Pk 50 ppm/ 10M/8H (SCP-U) FEREAC 39,23540,74. DOT: Flammable Liquid, Label: Flammable Liquid FEREAC 41,57018,76. Occupational Exposure to Benzene recm std: Air: CL 10 ppm/60M NTIS**. Currently Tested by NTP for Carcinogenesis by Standard Bioassay Protocol as of December 1980. "NIOSH Manual of Analytical Methods" VOL 1 127, VOL 3 S311. Reported in EPA TSCA Inventory, 1980. EPA TSCA 8E NO:12770027-Followup Sent as of April, 1979.

THR: Poisoning occurs most commonly through inhal of the vapor, though benzene can penetrate the skin, and poison in that way. Locally, benzene has a comparatively strong irr effect, producing erythema and burning, and, in more severe cases, edema and even blistering. Exposure to high conc of the vapor (3000 ppm or higher) may result from failure of equipment or spillage. Such exposure, while rare in industry, may result in acute poisoning, characterized by the narcotic action of benzene on the CNS. The anesthetic action of benzene is similar to that of other anesthetic gases, consisting of a preliminary stage of excitation followed by depression and, if exposure is continued, death through respiratory failure. The chronic, rather than the acute form, of benzene poisoning is important in industry. It is a recog leukemogen. There is no specific blood picture occurring in cases of chronic benzol poisoning. The bone marrow may be hypoplastic, normal, or hyperplastic, the changes reflected in the peripheral blood. Anemia, leucopenia, macrocytosis, reticulocytosis, thromocytopenia, high color index, and prolonged bleeding time may be present. Cases of myeloid leukemia have been reported. For the supervision of the worker, repeated blood examinations are necessary, including hemoglobin determinations, white and red cell counts and differential smears. Where a worker shows a progressive drop in either red or white cells, or where the white count remains below 5,000 per cu mm or the red count below 4.0 million per cu mm, on two successive monthly examinations, he should be immediately removed from exposure. Following absorption of benzene, elimination is chiefly through the lungs, when fresh air is breathed. The portion that is absorbed is oxidized, and the oxidation products are combined with sulfuric and glycuronic acids and eliminated in the urine. This may be used as a diagnostic sign. Benzene has a definite cumulative action, and exposure to relatively high conc is not serious from the point of view of causing damage to the blood-forming system, provided the exposure is not repeated. On the other hand,



URS REF 6

AN INTERNATIONAL PROFESSIONAL SERVICES 0994% 24

URS COMPANY, INC.

CONSULTING ENGINEERS

September 10, 1987

Mr. Russell Logalbo

Utica, New York 13503

Utica Board of Water Supply

570 DELAWARE AVENUE BUFFALO, NEW YORK 14202

TEL. (716) 883-5525

NEW YORK MONTVALE, NJ BUFFALO, ATLANTA AMPA HATO REY PR WASHINGTON DC 90STON CLEVELAND DENVER BALLAS SEATTLE SAN FRANCISCO SAN MATEO, CA

URS COMPANY

CEP 17 1987

JOB #_

Dear Mr. Logalbo:

Box 345

As I mentioned during our telephone conversation on September 8, 1987, URS Company, Inc. is currently conducting a Phase I investigation of the following sites within the City of Utica:

- Monarch Chemical Company, 37 Meadow Street

- N.Y. Emulsion Tar Products, Washington Street

- Mohawk Valley Oil Corporation, Lee Street

We are performing this investigation for the New York State Department of Environmental Conservation pursuant to the requirements of the New York State Superfund Law (Chapter 857 of the Laws of 1982).

This is to confirm our telephone conversation wherein you provided the following information:

- The source of water for the Utica Board of Water Supply is Hinkley Reservoir which is located approximately 16 miles northeast of the sites.
- The Utica Board of Water Supply is the sole water supplier for the area within 3 miles of the sites. This includes portions of the Villages of Yorkville, Whitesboro, New York Mills and New Hartford and the Towns of New Hartford, Whitestown, Marcy and Deerfield.

We would appreciate if you would review this information, note any necessary corrections, and return a signed and dated copy to indicate your concurrence. Your prompt attention to this would be appreciated, as the information is necessary to complete our evaluation of the site. Thank you for your cooperation.

Sincerely,

muyetta. mauche

Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC for Linda J. Clark, Project Geologist, URS Company, Inc.

MAM/bc 9/10/87L 35154/B3

I agree with the information as it is presented.

Mr. Russell Logalbo

Sept 14 1987





TOWN OF MARCY

P.O. Box 251 — 9455 Toby Road Marcy, New York 13403

November 12, 1987

URS Corporation 570 Delaware Avenue Buffalo, New York 14202-1207

Attention:

Muffett A. Mauche

Staff Engineer

Gentlemen:

 \quad Enclosed please find the information requested for your Phase I investigation.

Yours truly,

Karl P. Maxwell

Maje REN

Supervisor

 KPM:ccm Enclosure

> RECEIVED URS COMPANY

> > NOV 13 1987

JOB # _____

URS CORPORATION

570 DELAWARE AVENUE BUFFALO NEW YORK 14202-1207 (716) 883-5525

ATLATATA BOSTON BUFFALO BUFFALO CLEVELAND COLL'IBUS NEW. NORK PARAMUS NA SAN ANTONIO SAN FRANCISCO SAN MATEO SEATTE SEATTLE WASHINGTON D.C.

Mr. Karl Maxwell, Town Supervisor Box 338, Routh 291 Marcy, N.Y. 13403

Dear Mr. Maxwell:

September 15, 1987

As I mentioned during a telephone conversation with Ms. Cindy Cochi on September 15, 1987, URS Company, Inc. is currently conducting a Phase I investigation of the following sites within the City of Utica:

- Monarch Chemical Company, 37 Meadow Street
- N.Y. Emulsion Tar Products, Washington Street
- Mohawk Valley Oil Corporation, Lee Street

We are performing this investigation for the New York State Department of Environmental Conservation pursuant to the requirements of the New York State Superfund Law (Chapter 867 of the Laws of 1982).

In order to complete a Hazard Ranking System (HRS) evaluation, we request the following information for the portion of the 3-mile radius study area which lies within the Town of Marcy:

- Any community water systems (surface water or ground water, municipal or non-municipal) NO
- The number (or approximated number) of private wells used as a 0 sole source of water, or the number of buildings not on public 75-100 water.
- Is there a municipal system readily available to the above? 0 (Requires only a line to the street) year

The portion of the Town of Marcy which lies within the 3-mile radius study area has been located on a road map and the appropriate USGS 7.5 minute quadrangle. If you have any questions, please contact me. Your prompt attention to this would be appreciated as the information is necessary to complete our evaluation of the sites. Your assistance is greatly appreciated.

Sincerely,

muyetta. mauche

Muffett A. Mauche, Staff Engineer, LeRoy Callender PC for Linda J. Clark, Project Geologist, URS Company, Inc.

MAM/mb 35154





Soil Conservation Service RR#1, Box 126-C, Second St. Oriskany, N.Y. 13424

Muffett A. Mauche, Staff Engineer URS Company, Inc. 570 Delaware Avenue Buffalo, New York 14202-1207

September 14, 1987

Dear Ms. Mauche:

The following is information that you have requested for HRS evaluation on three sites in Oneida County.

- 1) N.Y. Emulsion Tar Products, Washington Street, City of Utica
- 2) Monarch Chemical Company, 37 Meadow Street, City of Utica
- 3) Mohawk Valley Oil Corporation, Lee Street, City of Utica
- * No irrigation of Agricultural crops within a 3 mile radius.
- * Agricultural land in producțion less than a mile None
- * Distance to Prime Land greater than 2 miles.

Sincerely,

Keben Mangine

Robin Mangini

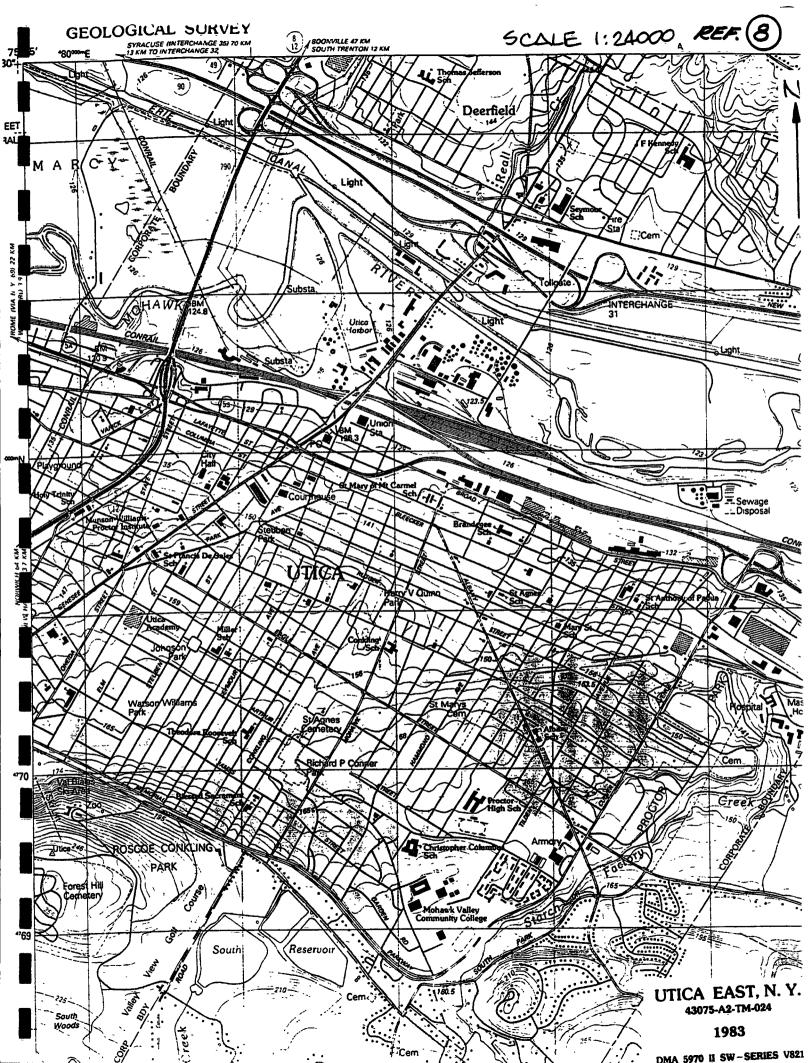
District Conservationist

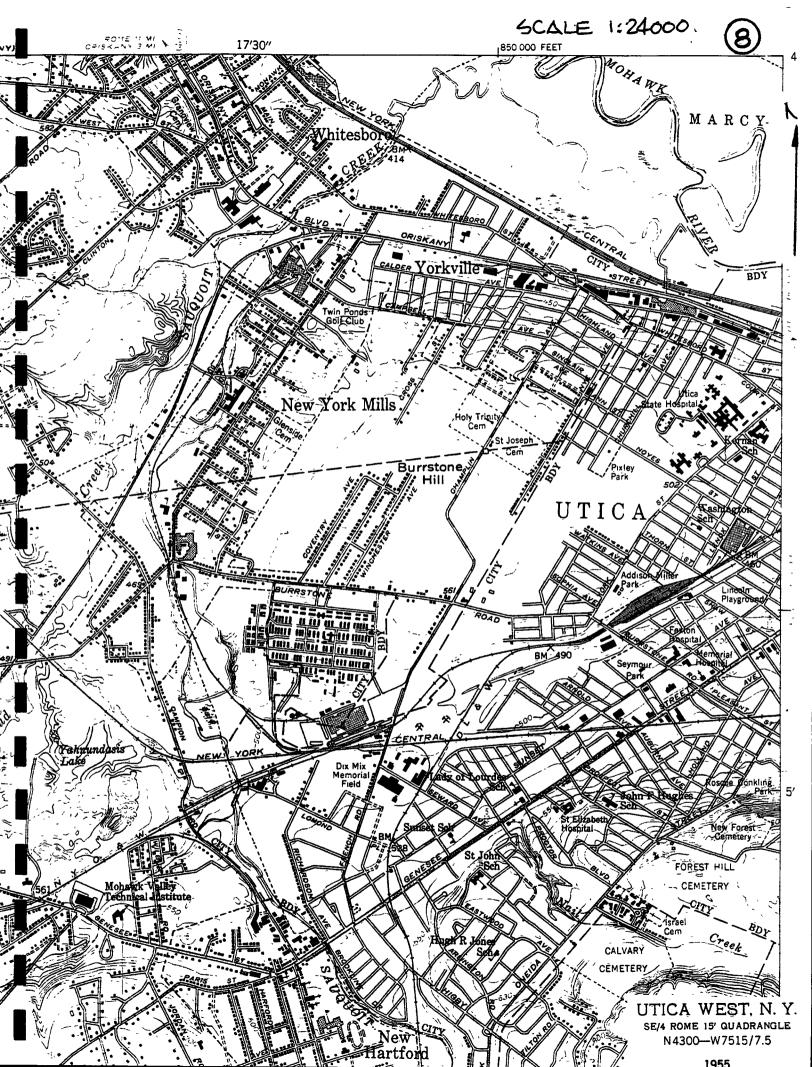
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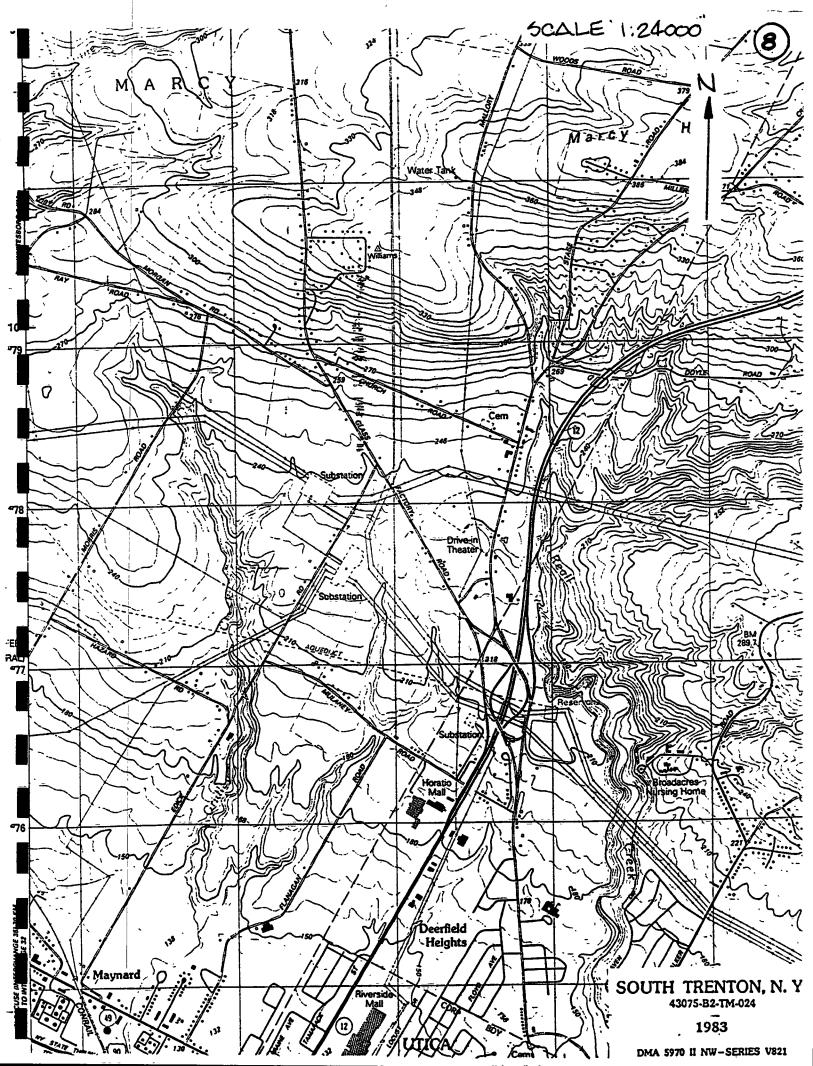
SEP 17 1987

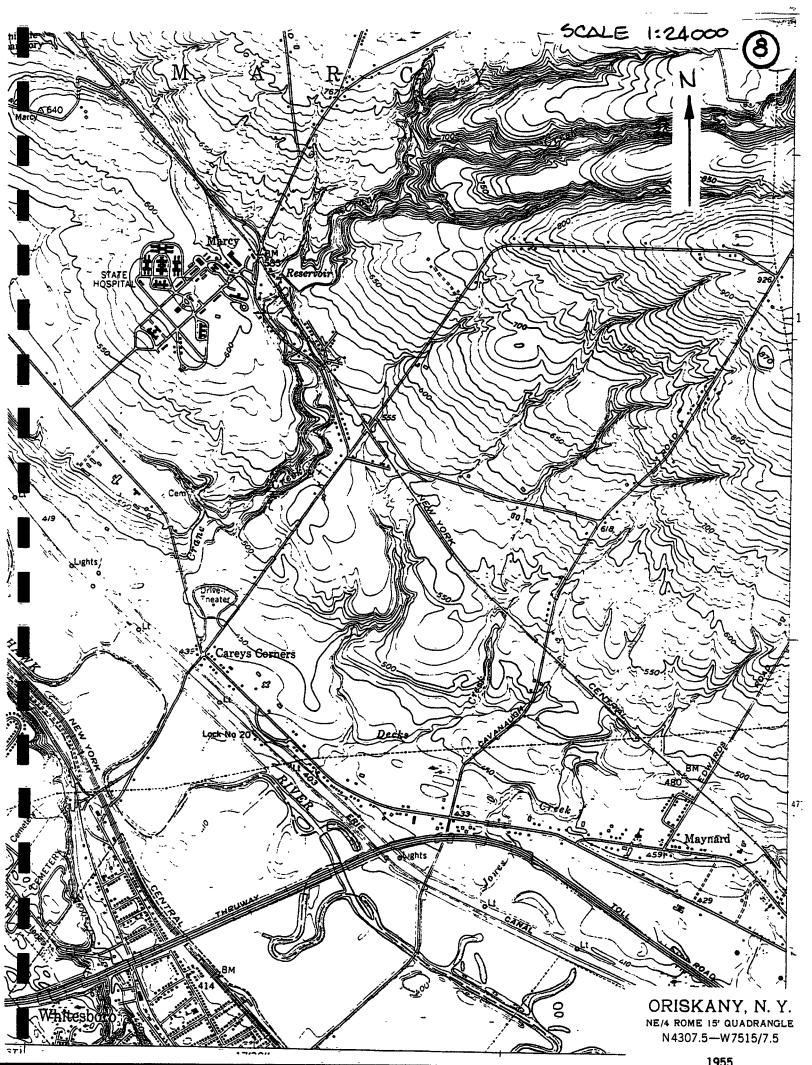
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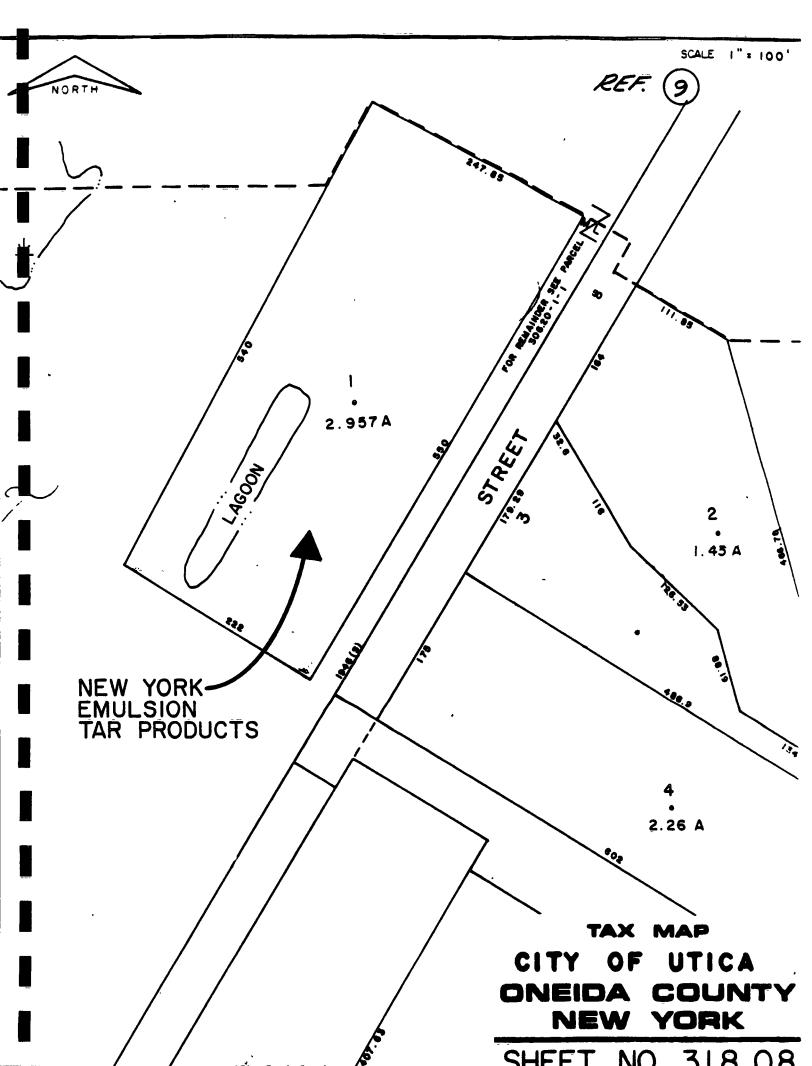




















Doing business as Central Asphalt, Cortland Asphalt Products, New York Emulsions, Northern Asphalt, and Western Bituminous Products.

> 1911 Lorings Crossing Road P.O. Box 5160 Cortland, N.Y. 13045-5160

RECEIVED

URS COMPANY September 16, 1987

SEP 21 1987

Dan Rothman,	P.E.
URS Company,	Înc.
570 Delaware	inc. JOB #
Buffalo, NY	14202

Re: Letter of August 18, 1987 (NYE-Phase I Investigation)

Dear Dan:

To confirm our phone conversation, Rod Birdsall and I have gone over the 21 points of information and made a few changes for clarification. Rod has talked with Bob Hannan, and we are sending a copy of this letter to him.

- (1) Koppers Company used to process material on-site. Tar Asphalt Service (the property across Washington Street which is now listed as being owned by Roselli Associates) would then pick up the tar material for use off-site.
- (2) Koppers operated on-site from approximately 1926 until March, 1977.
 Koppers was in the tar and asphalt emulsion business, at the site, at the time of our purchase. Suit-Kote is in the emulsified asphalt business. Suit-Kote purchased the site and inventory from the Koppers Company in March, 1977.
- (3) John Roselli purchased the Tar Asphalt Service property in 1977 and may be a possible source of information concerning on-site activities. This company may still be operating in the Utica area.
- (4) At the time of the purchase, Suit-Kote acquired the property and all inventory at the site, including asphalt and road tar. Suit-Kote operated the asphalt emulsion plant until 1983, at which time it ceased operation of the emulsion plant. Suit-Kote sold the road tar that was in inventory but did not process road tar at the site. Asphalt is the residue of the distillation of crude oil and is petroleum based. Road tar is the residue of the coking process of coal.
- (5) Some of the tanks on site had naphtha in them when Suit-Kote purchased the property from Koppers. Suit-Kote used the naphtha inventory in the production of asphalt emulsion and did not store naphtha after the inventory was processed.
- (6) There is lots of cleanup to be done on-site as of this time. The tanks are empty except for the residue in some of the tanks. However, there is some loose asbestos insulation on at least one of the tanks, and possible PCB oil in several of the on-site transformers.
- (7) Suit-Kote has a drawing of the site which shows the location of on-site tanks, etc., as well as the property dimensions.



Dan Rothman, P.E. September 16, 1987 Page 2

- (8) The on-site lagoon discharges to a manhole and then into the storm sewer system which is located on Washington Street. The lagoon was constructed by Koppers to comply with the spill prevention plan and countermeasure plan requirement for the facility.
- (9) There are a number of drums located throughout the site. Most of these are empty.
- (10) Suit-Kote produces no waste material in the emulsified asphalt production process. Everything goes into the product. Asphalt emulsions are a suspension of asphalt, emulsifiers, water and sometimes solvents.
- (11) A SPEDES permit is still maintained by Suit-Kote to cover the discharge from the lagoon into the storm sewer system. The SPEDES permit was originally applied for, and held by, Koppers.
- (12) In the emulsified asphalt production process, the basic input is a base asphalt, which is a very viscous, fairly solid material with a softening point of 120°+F.
- (13) The on-site laboratory still has material within it. It is believed that Koppers used this laboratory extensively. Suit-Kote used it only as a quality control laboratory performing very simple physical tests on the produced material, such as viscosity, which did not require any exotic chemicals. The contents of the old laboratory are unknown.
- (14) Suit-Kote emptied various tanks, after purchase of the property from Koppers, of inventory and used the material for various acceptable road building applications.
- (15) Suit-Kote used only a portion of the facilities during its period of operation on the property. Primarily, Suit-Kote used facilities located towards Washington Street for the production of asphalt emulsions.
- (16) The emulsified asphalt production process does not generate or use either benzene or naphthalene.
- (17) Jim Cromie was a long-time employee for Koppers and became Plant Superintendent for Suit-Kote after the purchase in March, 1977.

 Koppers Company has offices located in Pittsburgh. Jim Cromie retired from Suit-Kote in 1983, when we discontinued operation of the plant.
- (18) Rod Birdsall, President of Suit-Kote, has discussed the Utica location with Robert Hannan of Koppers. Mr. Hannan's office is in Pittsburgh. He is familiar with the site and former operation. Mr. Hannan can be reached at 412-227-2617.
- (19) Suit-Kote is aware of no spill or accident incidents during the period when Suit-Kote was operating the site.
- (20) During the period of Suit-Kote's operation, there were only one or two employees on-site.



Don Rothman, P.E. September 16, 1987 Page 3

(21) Again, Suit-Kote produced no waste products while operating on-site.

The above information is current and correct to the best of my knowledge. Please feel free to contact me if I can be of further assistance.

Sincerely,

Bill Fowlston

Bill fowlston

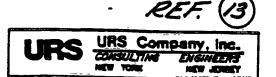
Director of Safety and Environment

 \mathtt{dr}

cc: Bob Hannan

HARBOR POINT PROPERTY LAND INVESTIGATIONS STEP 1 LAND REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York



Junes Dayle,
Sunitary Engineer

NYSDEC

207 Genesse St.
State Office Bldg.

Utica, NY 13501

JOB NO. 35154-01	
JOB TITLE NYS Superfund	
MEMO OF TELECON	
DATE 10/8/87 TELEPHONE \$ (315) 793-25	554
PERSON CALLING Muffett Mauche PERSON CALLED James D	oule.
REPRESENTING LeRoy Callender PC REPRESENTING NYSOEC-	Hi sa
PURPOSE OF TELECON AND/OR EQUIPMENT INVOLVED:	
Swuse- Mohawk River - Utica sites.	
TEXT OF TELECON	
Mohawk River is a Class C water wh	ick_
mean's fishing & secondary contact (boating)	
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@ Banny Brook [downstream] the R	vec
changes to class B.	
	and the same of th
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REF. (4)

New York State Department of Environmental Conservation

State Office Bldg., 317 Washington Street Watertown, NY 13601-3787 315-785-2261



Thomas C. Jorling Commissioner

August 18, 1987

RECEIVED URS COMPANY

AUG 20 1987

URS Company, Inc. Ms. Linda J. Clark 570 Delaware Avenue Buffalo, NY 14202

JOB#____

Dear Ms. Clark:

Enclosed is material requested by Ms. M. Mauche of LeRoy Callender, P.C. for your use in completing the Hazardous Ranking System evaluation for sites within D.E.C. Region 6.

Little Falls Landfill

a) No coastal wetlands within 2 miles

b) Freshwater Wetland LF-3 - see attached map

- c) No known critical habitats for endangered species nor national wildlife refuges within 1 mile. As critical habitat data is constantly upgraded, this should not be taken as a definitive statement regarding the presence of such habitats.
- d) No State Forest or Wildlife Reserve within 2 miles.

Monarch Chemical Company/ N.Y. Emulsion Tar Products/ Mohawk Valley Oil Corp.

a) No coastal wetlands within 2 miles

b) Freshwater Wetlands UE-2, UE-3, UE-4, UE-5, UE-6, UE-9 and UE-10 are within 1 mile of these sites. See attached map.

c) No known critical habitats for endangered species nor national wildlife refuges within 1 mile. As critical habitat data is constantly upgraded, this should not be taken as a definitive statement regarding the presence of such habitats.

d) Utica Marsh WMA is located in Freshwater Wetlands UE-2, UE-3 and UE-9. Izaak Walton League owns a sizeable parcel within UE-10 and/or UE-11 which would also qualify as a "reserve".

If you have any further questions, please do not hesitate to contact me.

Yours, truly,

Leonard E. Ollivett

Conservation Biologist II

Region 6

LEO: jes

Ence



NEW YORK STATE DEPARTMENT OF TRANSPORTATION

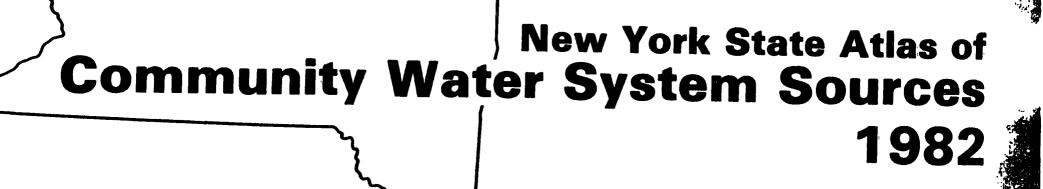




Freshwater Wetlands Classification Sheet for That Portion of ONEIDA COUNTY Outside The Adirondack Park -- 8/28/84

Map 31 of 38 Utica East Quadrangle

Wetlands Identification Code	Municipality	Classification
. UE-1	Utica	
UE-2	Utica	ĮĮ
UE-3	Utica	ΪΪ
UE-4	Utica	ΙΙ
UE-5	Utica	II
UE-6	Utica	II
ŰE-7	Utica	II
UE-9	Utica	ΙΙ
UE-10	Utica	II
UE-11	Utica	II
UE-12		II
UW-2	Utica	II
	Utica	II

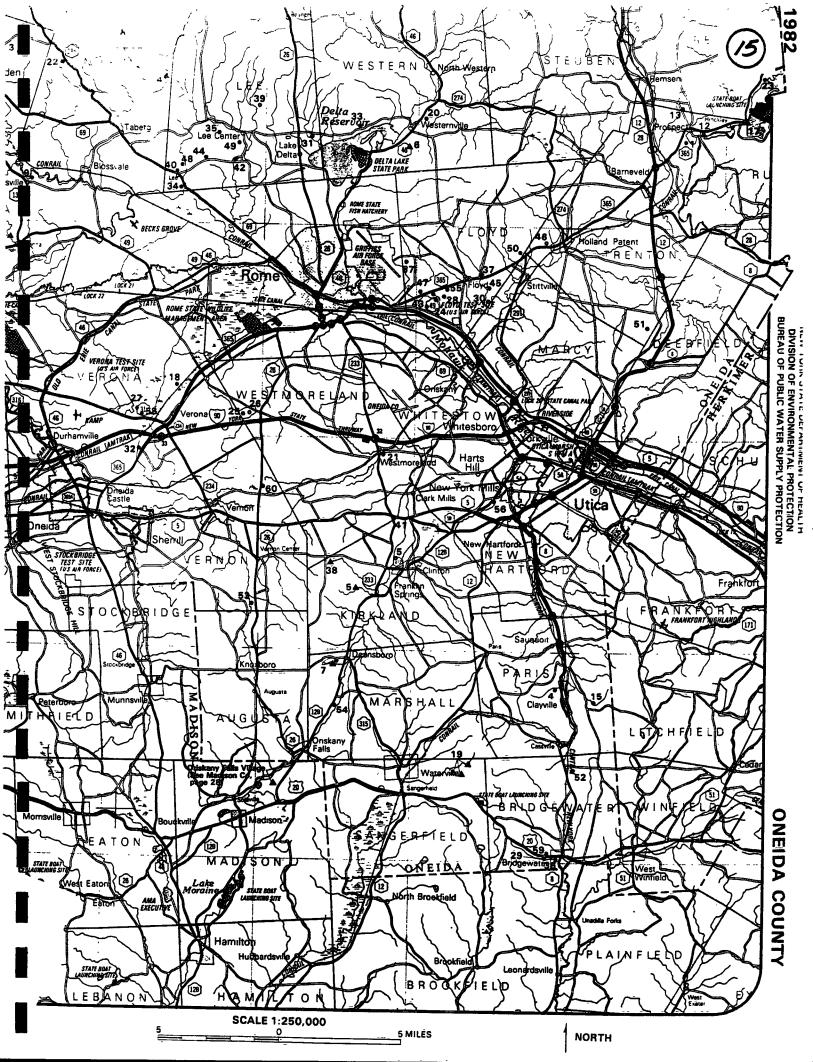


NEW YORK STATE DEPARTMENT OF HEALTH
DIVISION OF ENVIRONMENTAL PROTECTION
BUREAU OF PUBLIC WATER SUPPLY PROTECTION



ONEIDA COUNTY

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ID NO
   COMMUNITY WATER SYSTEM
                POPULATION
                     SOURCE
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Non-Municipal Community
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URS Company, Inc.

JOB NO. 35154
JOB TITLE NYS SUPERFUND
DATE 9-2-87 TELEPHONE 4/3/6/704-5/63
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REPRESENTING UPS CO. INC. REPRESENTING CITY OF UTICA
PURPOSE OF TELECON AND/OR EQUIPMENT INVOLVED: FIRE EXPLOSION
THREAT INFO, FOR HONAPCH CHENICAL CO
THREAT INFO, FOR HONAPCH CHENICAL CO., TEXT OF TELECON N.Y. EMULSION & MOHAWK VALLEY
HE SAID AS FAR AS HE KNOWS
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OF EXPLOSION THREAT", BUT AT THE
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WITH CHEMICALS SO "ANYTHING COILD
HAPPEL!"
CC:



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Number of Inhabitants

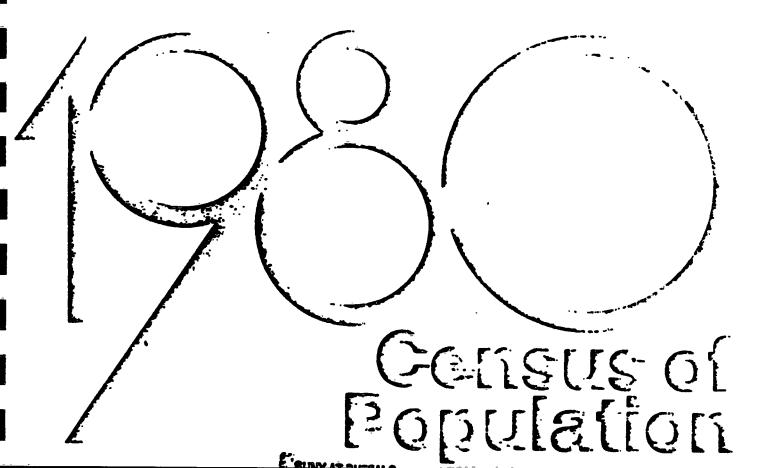


Table 5 Population of Places: 1960 to 1980

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			光器	3 785	Contraction with severe	College		1 410	1 426
Address ville		4 711	4 148	4 991	Company (\$100	Carrie	l	1 051 849	976
Balance Sen village	****	2 919	3 214	1 538 528	Contract Approves		•••••	2 001	2 239
Consessed (CP)		22	22	343	Outles viles			620	567
types of commences to the commences to t		14 703 L	17 336 1	18 210				677	656
1000 (IV		• 943	933	· 772	Outry Veloy village	(Italy)		684	66 1 }
Contraction (Contraction of the Contraction of the	*****	14 813		• • •	(100M ARM) 1000 1111			1 910	1 627
terter trade trade (QP) (Section	****	: 窓	8 232 11 119	•••	Caste 1889	Make	*****	399	3 605 1 065
Let Sun (SP)		10 / 	_ 1		0			1 300	1 332
Marana		.7 994	13 255	3 %2 13 92		Ul		1 412	1 200
beach Gly		17 22	18 233			*******		1 816	1 970
Constitution Laboration Contraction Contra		1 423 1		1 08				478 855	535 821
Infert (CF)		1 187	1 136	27	2 COluge Amile	Canally		5 636	5 771
hele ferry relief accessores helles accessores		18 106	18 43) 1	12 70 2 44	Charles seed (CD)			- 1	2 058
Man (Ch)		3 22	3 046	1 14				2 039	2 271
Action (Contract of the Contr		1 444	47	4		Ossil		1 193	i
And when a special of the second			1 918	N	(C)			2 491	2 829 1 4 368 1
Berger village	·······	1 095		i		MANUEL	******	5 272 902	897
bridge (CP)	******	14 140	18 555 2 509 1 276	! :	· (dept. 480)	TOURSESSESSESSESSESSESSESSESSESSESSESSESSES		18 144	16 653 '
by Ren (CP)	*****	? 때	1 元	1 :				402	-
to Pan (CP)		35 200	w 123	75 9				1 741	1 686 1
Bas Day 1809		14 840 2 972 702 55 880	44 123 1 367 3 910	1	(C) (C) (C) (C) (C)	*********	1	2 161	2 363
has bee will accessed the second		3 强		•				5 336	5 450
Will exceeded and the same of	*******	ä	554	1 4		0000000000 NAME 000000000000000000000000000000000000		8 869	8 70° 24 '38
Homesta viet				. 1	Calore village			34 719 7 123	5 728
Name (1884 1889) (1884)		1 151		. 1	Comment (CDP)		*******	330	347
Bearings-Helery Best (CP) Uster		1 002 9 308 1 345	1 2					1 254 2 342	7 401
Marin (D7)		1 345 2 344	7 🖛	6 2	400 Cassana (GP)	(tariff		656	734
Bearing Grow (CD)		1 140	1 19		··· (manufacture value) ··			20 132	.9 63.
Bronds vings		i 454 44 321	28 22	15	367 Capage (CF)	***********		24 752	1
		4 11	l					649	
7		2 47	1 12		Carto stage		*******	2 702 12 953	1 12 7
Charles (d. s)	******	1 %			The second second			3 164	
(man) (f) 60		1 1		38 1 1	714 Called 57			20 13	19 5.



Table 5. Population of Places: 1960 to 1980—Con.

	[k	A GARDS & A			ns since 1970 see table 4. For ma				
corporated Places passis Designated Places	Counties	1900	1970	1960	Incorporated Places Consus Designated Places	Counties	1900	1970	1940
	Seretoge	1 20	2 002	:::	feregreik vilage	Charlesse	904 509	908 562 1 733	905 453 3 737
pary times (CDF)	NOSS	2 786	2 377	2 040	fot láveti vilego	. Westington	3 561 646 1 376	711	874
1809	Grand	705	745	621	len Hangaray (CP)	Orange	2 555	2 800	2 809
	Westings -	4 447	7 333	9 815	for Pan vilus	Life	9 550 7 995	* ***	-
m Hagith (CDP)	Dutinos	3 225 1 737	3 292 1 735	1 949	frankler vilage	- Northway	2 995	3 305	3 677
Sales (CD)	Seffed	2 700 3 770	1 725		franklin village		27 051	352 32 156	525 32 403
Charles Agents	(2000)					4	1 887	1 948	2 124
	Longston Sufficial Sufficial Sufficial Sufficial Sufficial Controlled	30 394	3 436 32 274	5 440 14 726	Propinsi vilago	Contrages Contra	11 126	10 326	8 47
Aut (CP)	inflation	326	347	470	Propert village	Named	38 272	40 374	34 419 47
	. Scherochady	1 113	500	390 777	Immelest (CDF)		1 906	1 772	1 62
•		3 374	3 017	2 307	Municipia (CDP)	Migs)	13 312	14 003	1 23
(OA	. Alex	19 619	22 išš	13 580	Miles (IV	Mentalisty	777	812	81
W 1888	50				Generale vilege	Wysing	334 245	385 270	36 30
w ville	1000	1 897	2 061	2 025	GB147 1910			- 1	~
	Colours (pf Al	100	142	136	Geng Mills (CDP)		2 300 22 927) 258 25 373	23 94
	Suffet	16 542	24	19	Contain City village	***************************************	7 719 1	7 488	
		0.004	10 032				4 238 [4 614	
		1 053	10 050	1 007	General (CDF)	Manne	1 339	:::1	
(07)	Cathorn (pf 4)	1 053 26 073 10 053	io 333	1 260	(Crain) 1819	May	6 746	5 714	3 28
The same second		2 402	2 672	1 058	(ann) (f)		15 133	16 793	17 26
, , , , , , , , , , , , , , , , , , , 	Fullen (SI	125	175	185	WILL 07	Commo (47, 48)	15 133	16 793	17 26
		7 45	2 697	2 073	Character of the con-	Senam (# if)	455	552	52
Pais (09)	Putem	' 38 l	أفخة	47	Gia (07)	1867	1 179	1 149	
		1 761	! 40	263			24 418 2 632	25 770 2 720	23 6
M (Yells	1 554 15 310	1 539 16 855	i 25	Can Park William	Daletti	504	587	\$
# (#					Gos felt dly	Will assais seessais	15 927	17 222	18 5
(1)	160	765 363	1 090 377	1 004 370			4 754 17 836	19 477	21 7
	Contract (at at)	422	473	434	Galden t Bridge (CDP)	Walker	1 367	1 101	•
-	(10)	* 97	7 (93)	6 791		0	4 874	4 342	3 9
	(Marie	2 430	2 611			(**********************************	4 205	4 574	4 9
Congo Hamilto (GP)	(Militario)	20,385	23 790	•••	C	Second .	2 713	3 110	3 3
(D)		5 522	s ini	• • • • • • • • • • • • • • • • • • • •	***************************************	(Alternative Name of the contraction)	1 944	2 098	2 2
		i iii	1 753	1772		(d) (d)	312	1 012	1 0
		7 140 13 852	1 434	7 184	Grand Victorian Author		2 696	2 784	27
100 1000	5/10				Great Hart village	Carlomages (at 1t)	2 924	10 798 3 131	10 1
Manageret (CP)		13 967 39 317	15 926 46 270	14 779 44 036	Great Had Mark Village		5 604	6 043	4 9
		4 330	2 440	1 752	Green (CDP)		16 177	1 874	2 0
(D)		20 107	12 392	8 361	(COST) 1889	000000	2 696	3 297	3 5
Nachagas (CDF)	200	10 170	1 143		Granisus (CDF)		13 869	1 493	5 4
		. 45	134	94	1		2 273	2 481	2 6
Section of the consession	NEED	7 996 10 917	41 76	10 721	Grant War (GP)		1 571	• • • •	
Number Agets	Orentage	3 412	4 333	4 709	(Cont.) (CP)		955	2 002	2 2
	Namin	2 700 1 574	2 000	2 140	Granush viliago			2 262	1 2
				نندد	(THE 1888)			410	1 2
(QP)	Starting	3 600 541 730	2 %2 576	2 344	Harby Way	17	10 342	10 215	. 3
(101)	German	. 22	. 732	739	Hambur village		3 725	3 434	•
100		1 000	1 049	帰			1		1
		4 175	1 1	1 99	Hammanham' - Charles		1 045 7 256	1 066	i
grelle vilage		1 23	37	123	Hamping Park (CP)	SIG	1 331		
			1		House village assessment	(MAN)	1 526	1 688	1
(COMP (CDP)		122	374	4 317	Haraba villago concessorone	900		955	ì
(4)	Opini	4 期	4 706	5 157	Harman Sauth (CSP)	(100)	1 254 5 087	:::	l
un Hamilto Harin (CDF)	Onical	7 %	7 33	2 528 20 138	Harten village		. 44 1	. '	1
	Walder	1 1 14	1 2 911	3 795			. 937	834	1
eed (CD)	5/40	11 99	15 031 16 556	18 773	Nemate (GP)	Verice V	10 216	12 226	١.
NG (GP)		13 745	15 999	1	Control of the last of the las		20 940	13 957	•
dap	Malai	374	4	314	(37)		1 100	8 196	5
	MVW	45]	714		Northead (CD)	Andrew	3 576 5 010	3 447	1
		3	374	379 764				943	Į
	(1777)	13 415	15 317	۱	Harmonian Street			39 411	34
-	Marrie	5 970	6 474	5 507	Hartana villago	William	: ' ' ' ' ' ' '	521	'
			8 517 2 763	3 34	11		1	9 112	
		7 944 13 398	1 20	'			8 123	1 112	
	Wa	13 37	i sii	تقة أ	Harrings village announcement	sees S (gertina) essessessessessessessessessessessessess		770	H
•			I	1	(D)		6 994		
HATTONIA TANDO		4 701	1 446		Tables Sty Till Tales			1 512	1
		:	Si	1 52	Hardell Heat village		43 245		
Mark (GP)	(199)	44 45 55	4 02	2 62	1 (12) (12)	******	3 %	2 184	1
	 	: '##		قة أا	A lambour out after the course	(1949)	4 107	4 434	•
		14 60	19 44	17 47				ı	. [
		. 3	1 13	4 35	1 10 10 10 10 10 10 10 10 10 10 10 10 10	(1998)		1 05	
-			LI I∭Ž	1 🕮	4 10mm (DP)		5 732	,, , , , , ,	, ,

Table 5. Population of Places: 1960 to 1980—Con.

Her changes in boundaries of recommend places seen 1970 and highly 4. For making of symbols, and immediation

•		la tain a p	ovrderes ef =		1970 to 1980 4 W 1980	material (s) Marifelt (s) Season (s)			
Incorporated Places Consus Designated Places	Counties	1980	1970	1940	Incorporated Places Consus Designated Places	Counties	1980	1970	,,
* *******				İ			3 405	3 371	2 5
Hillis (CDP)	, Dydum	2 991	2 750	••••]	Lieve Harter village	Marine	24 444	25 390	20 4
Hallade Lake (CDP)	. Deldies		أففة و	1 336			9 670	11 626	11 5
Hallade Lobe (CDP)	. INCOME.	اذها	· 337	585			. 334	353	. 1
Hobert ville	Lifet	24 302			LONG BOOCH CITY	Neman	34 273 1 720	33 127 1 995	26 4
Helion village Helion village Helional (CDP) Helional village	. 0.000	952	778	440	Long Seach CDV	A	11 480	9 200	
Holand (CDF)	. [10	1 347	أغفة	538			3 364	3 671	3 6
Highend foton villige		1 1 1 1 1 1 1	1 868	1 786	Lineary (CP)	None	20 424	23 151	19 8
Highest (CDP)	Silled	13 515			Lynday (CDP)	. Oncretage	5 129	•••	
	•	3 425	4 143	3 422	Underste whose	Otem	914	688	•
Harrane falls village	. Cortigui	2 410	2 24	7 143	Lyan 1889	Wayne	4 160	4 496	4 0
Honorus Fells village	- American	3 (07)	3 (97)	4 023	Lygna fells willige		755	852	•
Hospital Junction (CDF)	_ Durchess	1 754 10 234	2 055	13 907	Macron when	Continue	1 184	1 319	1 1
Hornel City	· 20000 · · · · · · · · · · · · · · · ·	7 52	12 144	7 707	Makes view	Medair	396	386	. :
Horseld Junction (CDF)	. Ournett	7 348 3 081	2 753		Lyndy village	, Mineste accounts	7 668	5 265 8 048	ė
Housings (CDP)	Alegary	1 404 7 966	1 620	11 075	Materia of	No.	9 262	10 036	9:
Hartefoods Norm (CDP)	(chipita	7 419	8 940 7 917	7 752	Historia ville	- Nestille	17 616	18 909	17 e
Hudsen fells 18890	***************************************		' ''				1 498	1 305	٠.
Humay village	Gare		230	457	Manager VIII	. Omero	8 445	8 541	
Hymenstan (CDP)	5/14	2) 727	12 401	11 255	Harles All Consesses	Createst	5 241	4 295	1:
Huntagian Bay village		20 749	29 817	23 436			5 354	-5 488	
Marker (CDP)		4 072	4 061	. :::	Manager ville	Name	1 044	053	3 .
Hyde Park (CDP)	Declas	2 550 7 450	2 905	10 199		Complete	i 87 0	2 017	1
tion viliage		1 7 22	i Y 元	780	Marian Policy Village	. October	755	816	
ten vinge		1 229	143	10 362	Margania viliago		2 275	1 580	•
	Mat/40	57 646					7	. ~~	
		5 774	5 878	5 494	Manager (GP)	(1000)	24 454	26 421	32
incepton villago	 	4 047	5 3%	1 844	National Lag Apple and and an annual or an a		19 779	22 112 14 042	15
(D)		19.48	7 492	•••	Master (GD) Master (GD) Master Seath (GD) Materials (GD) Materials (GD)	Life	io 413 l		•
tels forest (CP)		3 70	24 224	29 799	Master Battle (CIP) annual contract	. Ma	8 318	4 870	3
		عتد ا	1	l	Marraces village		965 3 923	841 1 995	١,
	On the same and the same	35 775	39 795	41 018	Marmad (CIP)		7 511 1	272	'
Jamesiann West (CDP)		2 460	3 401	•••	Market 10077 *********************************	- Gran	2 007	1 534	1
isflered region (CDF)		13 300	فقة •		Majoris day	. Mile	944	981	Í
The same and a second second			1	۱	************	^	1 626	1 567	
Juffernande vilege Jeden (CPF) Jebest City vilege Jebesteen City		- 12 5%	14 010	10 74			5 500	6 247	•
(QF)		12 739	10 005	10 116	Medical (CP)		20 418		١.
		1 377	10 045	10 370	Hedra vilagi	Otas	6 392 2 171	6 415 2 2 189	, ,
inter vitae	(hardes)	. 1 371	1 473	1 390	Makes Pat (GP)		â i 19	6 661	•
	94		2 122	2 212	Manual villa	Contactor	4 012	3 449	Z
Konsult vilago	Chairm (at at)	. 1 055	1 210	1 231	Maridon	(M)	344 24 478	369 25 904	10
	Course (six set	970	112		Maria (CD7)	Names	3 661	2 564	
Common village	(10	10 474							
Catalana (CD)		i ia	1 243		Manage village	Company	1 621	1 555 1 410	1 :
Coule Park (CDP)	··· Yell	1 15		نفنا أ			3 227	2 327	
Company and		*1 4 19	1 233 5 555	1 071	Marie Ward (CD)		5 703		l .
Company (CDF)		1 2	5 614	5 410	Making		1 995 21 454	2 132 22 607	23
Granton City	Mills	24 40	25 544	29 24	Middleban dy		" 🚟	725	
Level 100 village		7 000	'I ···			August	514	527	1 .
Lockson COV		22 701	28 457	7 55	Military	Dutos	7 377	1 735	' '
(acone video)	(1118)		11 336	33		10100	, , , ,,	•••	
Later (CDP)	9.00	- 210		7 79		Determ	1 013	1 042	
Lake Carmel (OP)			1 4 47	3 1	Mil Nest 1889	100000	959	982 480	
Latin George village	Water	1 96	/ 1 9 #	102	The second second		2 043	1 861	
Late Group village	··· Mile ·····	2 01		1 1 14			20 757		
AND ENGINEER PLANTS OF THE PARTY OF THE PART				1			20 757	21 845	x
Lake Lyzanne-Hadley (CDF)	1000	1 199		2		(m)	925	1 967	'
_	Souther (F. 11)	_ 176			Mind The assessment			7 245	
Late Floral village		_	2 73	i] 2 🙀			7 756	3 301	1 .
Links Backerberry (CT)		2 48 2 19 2 19 2 19	1		حصيدان	Car	5 994	4 439	4 :
		- { ?		2 9	Marie Sentent (DR)		1 267	l	.
(Abouted William		1 14	11 3 200	4 3 77	3 Mariani (COM) aireann ann ann ann ann ann ann ann ann ann		1 1	8 797	1
		13 05	13 34	5 12 25	6 Marie (GP)		2 316	1 53	3
(Clare (Sept	[614]	- 19	7 💥	اخا أغ			6 306	5 99	1 1
Latina (CP)	Waite T		31 j#		Marine Mile Will accessor		1 791	1 53	
			_	1					
Granden Alega	1000)	!! 12			Martin ville	Capage		67	
	(*********************************		Å 22	ي اه	NA 1		1		. 1
CONTROL VICENT AND ADDRESS OF THE PARTY OF T	Viigo		N 4 54	āl se		Carren	- 721		
(ACCOUNT			2 34 0 5 11		Merricon vingo	S Gureta	2 757	2 29	
is lay viley	(anim)	57 0			74 Hunton Late (CP)	Malan	1 730)	1
		" " "	b 121	2 3 3	20 Haut Cats -Bays	(ringe	6 025		
Littly village	Mager		73 16 51	4 7				1 (1
LING VIEGO	with	2 Œ	5 14	b 13		West	46 71	72 77	
ion introduction (CS)	(Artical		n						
Gresiere view	(#17200	4	ا 🖮	15 1	39 Number Park village		. 2 60	- I	i
Larrie Park (CDP)		1 24	2 8			(*********************************	. 2 47		
Landarduria vallego	1744	75 ¥		20 3	35 Manustan vilia	(5100)	2 72		
	Mandagemen	1	56 7 6	 	35 Myers Carner (CSP)	Name of the second	5 18 12 57		
Liftle Velley village		! ?			44 Numer (GP)		1 20	٥l	i
		4 9		2 3			. ' 44	5 1 3	
			# 1 i #	# i		Agrander	1 26	5 1 4	

Table 5. Population of Places: 1960 to 1980—Con.

		(for designs of	-	iconteción l	facos senco 1970, seo table 4. For m	sonny of symbols, use extraduction)			
incorporated Places Consus Designated Places	Countles	1980	1976	1960	Incorporated Places Census Designated Places	Counties	1980	1970	1960
tensis late (CP) takene viles takene viles tenses (CP) tenses (CP) tenses viles	Manganay	1 304 491 547 10 704 10 017 1 190 2 436 2 436	716 583 10 046 11 444 1 266 1 349 26 219	729 555 1 964 12 866 1 234 1 762 30 979	Oristor Bay Core village Posted Poor village Posted Poor village Posted Village	Nation Studio Management Conference Conference Safeth Conference Confer	1 709 2 196 404 3 729 511 535 11 291 1 996	1 320 2 494 401 3 774 489 634 11 582 1 914	988 2 570 578 3 476 450 567 6 838 1 734
New City (CIP)	Rectard	7 435 35 459 3 120 1 532 1 365 2 313 7 401 4 736 744 70 794	8 721 27 344 2 556 1 111 2 453 10 114 4 056 908	1 423 1 423 10 808 10 808 3 061 827	Peach Lake (CDP)	Portid	1 464 998 446 15 973	17 146 19 283 2 076 6 673 5 293 4 538	18 737 1 964 6 114 5 770 4 629
top fair (lap)		70 794 1 750 7 612 2 120	75 385 1 156 8 883	76 812 4 041	An (C)		405	433 1 261 1 989 858	1 887
	Total	7 01 97 77 77 77 77 77 77 77 77 77 77 77 77	1 539 233 1907174 275 443 3 605	7 781 184 1 424 815 2 427 319 1 479 281 1 409 578 221 991 3 788 102 394	Profes village	- Chindre - Chin	- 557 2 357 2 369 - 216 - 216 - 383 - 6 83	1 674 2 617 2 384 373 1 183 247	868 1 750 2 408 1 906 345 1 076 180
tage Yest ANDs village		13389289	65 613 636 6 186 1 120 1 379 11 726	162 394 643 312 1 353	Amate (GF) Amate (GF) Amate (GF) Amate (GF)		- 9 45 55 55 55 55 55 55 55 55 55 55 55 55	1 755 10 759 31 495 1 593 1 082 835 18 715	1 749 21 973 27 710 1 379 1 025 705 20 172
terris Balletin (GP)		19 019 1 350 35 020 20 430 7 432 2 743 1 464	29 526 1 276 22 892 5 903 1 425 1 475 12 000	19 439 1 574	Retitions War (GP) Research other Research other Research (GP) Research other Research ot	Distant	5 905 1 210 1 253 6 749 1 031 553 2 421	7 078 1 372 7 110 	5 877 564 295
Remin Haller village Remin Haller village Remin Handel village Romin Haller Fide (CDP) Romin LanderScraft (CDP) Romin Haller Fide (CDP)		786 1 567 2 307 11 511 21 305 12 640 15 114	910 910 11 117 29 123 13 450 18 154	450 337 917 12 974	Part Heavy ellips		1 450	1 330 25 803 2 132 2 602 1 532 5 515 7 403	1 201 24 960 2 295 2 622 1 767
North Terro Ryde Perk (CDP)	Warring	15 114 7 126 7 451 1 171 7 970 7 994 35 740	18 154 5 232 7 484 8 467 8 334 34 012	17 929 5 972 7 412 8 818 34 757	For Jufferson village	Contraction	8 499 740 1 136 14 521 3 147 10 435 29 757	8 852 842 1 304 15 923 12 883 10 303 32 029	9 268 898 1 336 15 657 722 7 765 38 330
Restricts (GP)		14 530 - 364 - 2 550 12 457 - 459 - 467 - 2 457	14 481 1 192 15 653 6 663 2 698	1 156 9 175 2 200	Marki ologo	Correspondence	2 415 946 1 398 1 401 3 091 645	2 480 865 1 498 3 034 2 797 626	2 254 692 1 414 2 410
Numb village		1 189 4 448 4 600 1 791 135 23 487	1 254 6 450 7 334 1 964 169 35 372	1 234 6 642 2 676 111 20 466	to Oak and IGM	Delication	1 692 5 236 621 1 868 9 047 340 2 542	1 660 3 919 602 10 136 332 2 336	1 719 567 10 506 375 2 093
Copin (CDF)		413 12 375 1 571 6 215 1 574 629	14 554 1 592 7 004 1 705 012	573 14 122 1 215 1 126 373	(2)	P (Arriva)	494 1 561 792 336 8 977 1 205 6 339	462 1 540 826 334 1 173 7 565	403 1 630 743 292 1 247 5 830
Character of Chara		1 061 3 277 16 287 10 610 751 14 933 5 120 3 671	2 667 19 169 11 658 700 16 030 4 340 3 722	2 044 21 046 11 477 754 13 412 3 270	Andreille Courty village	States	684 5 400 1 625 241 741 25 412 7 012 11 465	911 1 849 295 011 27 444 5 351	1 030 1 900 318 611 26 355
Orekery village		1 480 802 20 194 19 773 1 087	1 427 927 21 489 20 913	1 580 977 10 442 22 155 975	Section (CP)		43 826 42 205 14 109 1 136 2 134	50 148 47 455 15 008 -	51 646 12 883 2 681
Coup day		753 444 4 344 1 745	956 923 779 5 152 1 946	894 769 5 417 1 871	Batha Hatter officer accommons	Manual descriptions of the second of the sec	272 127 4 544 22 933	1 420 1 125 7 242 25 214	1 289 925 16 871

Table 5. Population of Places: 1960 to 1980—Con.

Incorporated Places Consus Designated Places	Counties	1980	1970	1960	incorporated Places Consus Designated Places	Countles	1980	1970	
Round Labe village		791 2 766 1 059	2 250	2 ié	Signy Peer (CDF) Sepredo (CDF) Sifteri value	- facilité	8 484 1 387 10 794 1 203	8 270 1 106 8 273	;
Regisally velocity	Total	548 148	546 194	445 159	Sylvan Booth village System (IDP)	Country Orange Orange Orange Orange Orange Orange Orange	1 243 9 818 170 105	10 084	216
head Garden vilage	Yeres (at all	1 263	1 207	1 156	1000 1000 1000 1000 1000 1000 1000 100	Rediend	8 267	7 424	
No are		15 083	15 860	4 游	Teryteen was	.,	10 340	11 115	I.
Saide feet vilege	A Parish	2 561	2 343	2 346	Therese village	Jefferen	2 664	7 811	
St. Generative (CP) St. James (CP)	. Ma	2 587 12 122 1 974	10 🕮	3 524 2 196	Remoted (CDA	Position	7 197 2 938	4 674 3 266	
Si jakopallo vililija Salamanto GTV	Managemery	990	2 007 7 677	1 449	Table 1007	- Vietra	1 529	1 256 739	
		950	1 025	1 074 29	Temperate div		18 693 72 795	21 898	Z
Series village	Westerspiele	1 293 2 742	1 107	2 161	Touri Law (CSP)	to	2 917	2 43a 1 18a	
Sandy Print village	- Canada	745	2 916 731	407			54 438	62 918	
Sprease Lake village		5 578 1 442	4 004 1 445	4 421 1 780	Transpirery village	Control Contro	7722 6 076	1 803 6 236	
	fema (st st)	4 116 23 706	10 20	4 441	Total International		1 049 4 478	4 654	
Sanjirina villaja	Library	3 912	4 190	16 430 4 286	Top (Cap		294 807	293 861	
		932	11 20	104	Lincoln (Con	Otalia	1 367 20 016	1 489 22 077	2
Sorrier (GP)	Washing	17 436 477	19 227	17 144			1 201	1 163	
School of annuaries	- (derected)	U 972	77 798	720 81 662 463	Digner des	Control	574 1 245	\$76 1 182	
Schröde village	(778)	1 014	1 125	1 146	Upper Name willige	Audiend	1 906 75 422	2 096 91 373	к
Scholary village		7 352	2 119	1 361	THE REP LAND SECRETARION		3 194	1 286	
Sprogram (CDF	as (right accessors	7 220	7 370	7 625	I Admit Charles Street Street		8 214 554 25 749	4 007 481	
Sprinds vilip	Maisti	5 344	5 070	الله ذ	Valley Sites will		35 749 559	40 413 522	
Salut (CDF)	Nesti	17 259 1	17 379 11 613 7 784	14 719			1 373	1 100	
Spring Fells village	Mil	7 444	7 794	7 409	Wine (2)	Oneith.	1 057 2 170	2 i 87	
		514	421	351	Victory village		571 1 707	718 1 475	
States was (GP)	Scholo	1 115 1 561	ı diğ	ففة ١	Valo (GP)		5 340 3 320	\$ 136 2 826	1
Summer village		2 630 2 630	709 2 706	2 92			980 5 459	955 5 277	
Surrected ICE (CDP)	SAL	7 344 18 072	% iš7	:::	Well (D)	Change	2 064	1 849	
Sharakan village	Suffish	555 1 449 4 661	\$34 1 516	1 36 1 36 5 15	Walter village	Disease	3 329 1 475	3 744	
Salary village	D dd 100	4 061	4 769		Water Falt (GP)	Grange	549	586 21 873	1
Shor (rest village		3 000 801	3 場	3 310	Mahada (ap	DAMES	5 110 1 818	5 607 2 017	
Section to the section of the sectio	(1000000	2 77	1 1 665	1 2 72	Wagneyer Feb. Sar (CP)	Dispin	1 799 2 834	2 743	
Sam village		A 550	5 276 3 134	5 00	A Marian Maria Constitution		4 120	3 610	1
Southern (CP)	*** ***********************************	3 194 30 106 225	فنذ	ä	Miletik tillige annansassassassassassassassassassassassa	**** **********************************			1
Some village	Worth	· 元	1 613 1 772	1 64	ودورون والمراكب والمتحدث والما		2 第	204 1 887 2 879	1
	A	7 140	. 200	9.73	a lateratura de la constanti d	(411)	2 405 3 303 27 661	5 418 30 787	
Sound Bossin (CDP)	1414	4 000	4 10	4 50	والمستعدد والمستعد والمستعدد والمستع		1 672	1 100	1
South Commandate	Sain	1 195	1 414		حطاحه منفض ا		2 440	2 716 5 26	1
South Dayless village	5470	16 437	1 99 20 44	. 14 31	O Waisely white accommons	1000	1 64	2 62	
South Personality (CPT		は常	1 000 4 011			Harri	S 499		
Sain HE (CDP)	,,	\$ 276		· ··	Westpart vilege	(1741)	64 5 76	77	•
South Hudson Falls (CDP)		1 955	2 007		With the contract (SP)		62	6 42	4
Sein Lader (GP)		湯端	1 34			Mil	41 69	• •	
Some (CSF)		8 199	200				. 1 824	4 2 04	7
- Court Velor Server (CDF)		1 342	10 57			Oarwy	5 44		ı
Spinished (QP)	Notes	4 14		•	West feet (CDF)	(high	171	3 65	11
	••••			1 .	West Gloss Fells (CDF)		1 13	4 1 15	
States agel	Nestin		i is	4 7	47 Westergen Bend -Cop	Contraction Contra	1 62	11 6 55	8
Serena (CDA		. 1 197		. 1			10 55	11	15
Studien spile	(m)	20 S2	4 35	5 10			7 M	12	74
Similar (CD)	(10	2 20	1 21	6 11				1 6 3	
Statute (GP)	Here	1 02/			Harris (GP)		3	53 5 5	٥
Course (CDA	Orenen	2 707			100 and 100 an	Other	1	05 13 6	73
Street William Ages	****	1 13			22 1100000 10000 000000000000000000000		· 1	\$3 1 0	75 i 86 i
Street Main Age		1 13	1 4		22 1100000 10000 000000000000000000000		· 1	53	



Table 5. Population of Places: 1960 to 1980—Con.

[for changes a basedories of accompanied places unto 1970 and high 4. for manner of accompanies

	(10.000)	. constant in	-comment t	PARTY SPECT 1970 Set table 4	No second of seconds	
Incorporated Pieces Consus Designated Countles Places	1900	1970	1960			
West Series (CD7)	5) 210 6 169 979 3 241 46 990 4 490 1 093 1 740 6 017 8 216 1 135 1 496 1 135 1 7 043 1 128 1 12	7 253 1018 3 764 50 346 4 805 1 955 1 961 6 878 9 154 1 264 1 671 817 1 671 1 671 1 671 1 773 1 773 1 7 716 1 7 716 1 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	940 4 014 50 495 4 794 1 049 1 300 6 314 8 295 1 320 1 026 1 441 1 034 907 907 907 907 907 907 907 907 907 907			



Site Inspection Report

OFDA	PO	TENTIAL HAZAR	DOUS	WASTE SITE		I. IDENTI	FICATION	
 €EPA	PART 1 - SITE LOCATION AND INSPECTION INCOMATION						02 SITE NUM	18EA
N CITE NAME AUD LOC		E LOCATION AND	INSPE	CHON INFORM	MATION	NYSDEC	#_6330	031
II. SITE NAME AND LOC			log STREE	T, ROUTE NO , OR S	DECIENC LOCATION	10544919		
New York Em	ulsion Tar Prod	ncts		shington		ICEN I INEK		
03 CITY		,		05 ZIP CODE	Toe COUNTY		Tozco	UNTY DE CONC
Utica			NY	13501	Oneida	ı	8	DE DIST
09 COORDINATES	1 LONGITUDE	10 TYPE OF OWNERSH			1			
430 06 37"	750 13 38"	② A. PRIVATE □ F. OTHER _	U B. PE	ZERAL	C. STATE	D COUNTY G. UNKNOW	🖸 E. MUN M	ICIPAL
III. INSPECTION INFORM 01 DATE OF INSPECTION	MATION 02 SITE STATUS	03 YEARS OF OPERAT	2001					
8, 6,87	□ ACTIVE	_ 19		1 1983	,	Marian Marian		
MONTH DAY YEAR OF AGENCY PERFORMING INS	INACTIVE		NNING YEA			UNKNOWN		
☐ A. EPA ☐ B. EPA C								
□ E. STATE Ø F. STATE	CONTRACTOR URS CO	rporation	☐ G. OT	inicipal 🗆 d. a Ĥer		MCTOR	(Mame di	lum)
05 CHIEF INSPECTOR		OS TITLE		7.77	(Seecify)	TICAL	OS TELEPH	ONE NO
Daniel W. Rotl	hman	Project I	Manage	er	URS Cor		1	8 83- 552
09 OTHER INSPECTORS		10 TITLE			11 ORGANIZA		12 TELEPH	
Gregg Townsend	đ				NYSDEC	Reg.6		793-255
				-			()	
	1			<u> </u>	37.2	· · · · · · · · · · · · · · · · · · ·	()	
						~~~~		
							()	
13 SITE REPRESENTATIVES IN							()	
		14 TITLE Safe	ty 1	SOX 5160	Lorings C	rossin		
William Fowlsto	on	Director						753-608
-				Cortland,	New York 1	.30 45- 5: -	()	•
							()	,
						· · · · · · · · · · · · · · · · · · ·	()	
•				- 12 - 12 - 1			()	
<u> </u>			-	<u> </u>				
							()	
17 ACCESS GAINED BY	18 TIME OF INSPECTION	19 WEATHER CONDI	DOMS					
CAPERMISSION WARRANT	11 A.M.			Sunny, 75	o _F			
IV. INFORMATION AVAIL	ABLE FROM	- 	·	· · · · · · · · · · · · · · · · · · ·				
01 CONTACT		02 OF (Agency/Organiza	EDR)		-	Io	3 TELEPHON	E NO
Daniel W. Ro		URS Corpo	oratio	'n] ;	716 ,88:	3-5525
04 PERSON RESPONSIBLE FO	R SITE INSPECTION FORM	05 AGENCY	06 ORGA	NIZATION	07 TELEPHONE N	a lo	8 DATE	
Kevin H. Sie	pel		Sá	ame	same		11 /1	11,87
EPA FORM 2070-13 (7-81)								

\ /	

POTENTIAL HAZARDOUS WASTE SITE

L IDENTIFICATION

♥EPA			SITE INSPECTION REPORT PART 2 - WASTE INFORMATION				NY CONTRACTOR	NJEMUF
				EIRFORMATIO	n		VYSDEC #63	3031
	TATES, QUANTITIES, AN							-
01 PHYSICAL S	TATES (Chock of the apply)	02 WASTE QUANTI	f waste quantities	l .	TERISTICS (Check of that a			
□ A SOLID □ E SLURRY □ 8 POWDER, FINES □ F LIQUID □ C SLUDGĒ □ G. GAS □ D. OTHER		TONSunknown		Ø A. TOXIC		CTIOUS		SIVE IVE
		NO. OF DRUMS	COME TARGET				M NOT AF	
IIL WASTE T	YPE	<u> </u>						
CATEGORY	SUBSTANCE N	IAME	01 GROSS AMOUNT	02 UNIT OF MEASUR	E 03 COMMENTS			<u> </u>
SLU	SLUDGE		1	100 0.00.00.00.00.00.00.00.00.00.00.00.00.0	E 04 00 mmeruit			
OLW	OILY WASTE		TARREST AND					
SOL	SOLVENTS			 	 	-		
PSO	PESTICIDES			 	-			
occ	OTHER ORGANIC CHEMICALS		unknown		 			
ЮС	INORGANIC CHEMICALS				†			
ACD	ACIOS				†		 -	
BAS	BASES				+		4 14 14 14	
MES	HEAVY METALS				 			
IV. HAZARD	OUS SUBSTANCES (See Ad	poends for most frequent	ly cand CAS Mumbers)	<u> </u>				
01 CATEGORY	02 SUBSTANCE NAME		03 CAS NUMBER	04 STORAGE/DISPOSAL METHOD 05 CONCE			CONCENTRATION	08 MEASURE OF CONCENTRATION
OCC	Naphthalene		91-20-3	ground sur	ground surface		,300	ma/ka soi:
OCC	Benzene		71-43-2	ground sur	face	-	5.40	mg/l GW
						—		
						 		
						t		
						\vdash		
				 		 		
						 		
				T		 		_
							=======================================	
								_
•	-							
V. FEEDSTO	CKS (See Appendix for CAS Number	ore)		<u> </u>	·····		 	L <u></u>
CATEGORY	01 FEEDSTOCK NAME		02 CAS NUMBER	CATEGORY	CATEGORY 01 FEEDSTOCK NAME		ME	02 CAS NUMBER
FDS				FD8		-		
FDS	·		<u> </u>	FDS				
FDS			 	FDS				
FDS			1	FDS		·········		
VI. SOURCES OF INFORMATION (Cre apacific reforances, e.g., 1000 files, sample analysis, reports)								
			.					
	Harbor Poi Niagara Mo		y Investiga	tions - Lar	nd Reports -	. 19	84-1986-	

POTENTIAL HAZARDOUS WASTE SITE

I. IDENTIFICATION

SITE IN	SPECTION REPORT	!		2 SITE NUMBER
PART 3 - DESCRIPTION OF H	AZARDOUS CONDITIONS AND INCID	DENTS	NYCEPEC	*******
II. HAZARDOUS CONDITIONS AND INCIDENTS			NYSDEC	#633031
01 Ø A. GROUNDWATER CONTAMINATION	02 CXOBSERVED (DATE: 1984 inves	** ~~ +r		
03 POPULATION POTENTIALLY AFFECTED: 456	04 NARRATIVE DESCRIPTION	Elgause,	DENTIAL	ALLEGED
A number of contaminants detected indicate site is source.		centrat:	ion line	es
01 🗆 B SURFACE WATER CONTAMINATION				
03 POPULATION POTENTIALLY AFFECTED:	02 (1) OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION) [] PC	POTENTIAL	□ ALLEGED
None reported or believed likely.	,			
01 □ C CONTAMINATION OF AIR	02 C OBSERVED (DATE:)	, <u> </u>	OTENTIAL	7 11 2000
03 POPULATION POTENTIALLY AFFECTED:	04 NARRATIVE DESCRIPTION	, ,,,	TENTIAL	C ALLEGED
None reported				
-				
01 C D FIRE EXPLOSIVE CONDITIONS				
03 POPULATION POTENTIALLY AFFECTED:	02 (1 OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION) 0 PO	OTENTIAL	O ALLEGED .
None reported				
01 & E. DIRECT CONTACT 18 909	02 OBSERVED (DATE:	- V P		
03 POPULATION POTENTIALLY AFFECTED: 18,908	04 NARRATIVE DESCRIPTION	AJ PU	OTENTIAL	ALLEGED
The site is only partially fenced				
**** *********************************				
01 & F CONTAMINATION OF SOIL ADDITION 30				
03 AREA POTENTIALLY AFFECTED: APPLOX 30	02 OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION	X0 PO	TENTAL	□ ALLEGED .
(Acres)				
Northern part of approx. 64 acre i	usrhor Doint property not	- conejí	¹A	
threatened or affected	ignoot some brobers we	COURT	.erea	,
OI TYG SOMEWING WATER CONTACTOR				
	02 OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION	★□ PO T	TENTIAL	□ ALLEGED
Approx. 120 wells within 3 mile r				
approx. ISA monto andmin a manage	adius, Deergleid + marcy			
01 A WORKER EXPOSURE/INJURY	02 OGSERVED (DATE:		<u></u>	
02 WOOVERS SOTTO THE CONTRACTOR	02 LI OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION	CI POT	TENTIAL	□ ALLEGED
None reported				
·····				
01 St. POPULATION EXPOSURE/INJURY				
	02 OBSERVED (DATE:) 04 NARRATIVE DESCRIPTION	₽ POT	TENTIAL	C ALLEGED
Sum of population potentially exp	to direct contact			e e e e e
subject to contaminated groundwate	OSEG TO GILECT CONTROL, .	and the	se pote	ntially
				

POTENTIAL HAZARDOUS WASTE SITE L IDENTIFICATION SITE INSPECTION REPORT O1 STATE 02 SITE NUMBER PART 3 - DESCRIPTION OF HAZARDOUS CONDITIONS AND INCIDENTS NY MYSDEC 63303I II. HAZARDOUS CONDITIONS AND INCIDENTS (Compued) 01 [] J DAMAGE TO FLORA 02 C OBSERVED (DATE: ____ **POTENTIAL 04 NARRATIVE DESCRIPTION** None reported 01 G K DAMAGE TO FAUNA
04 NARRATIVE DESCRIPTION (MICHAGE AN 02 GOBSERVED (DATE: __ POTENTIAL. None reported 01 E L. CONTAMINATION OF FOOD CHAIN 02 OBSERVED (DATE. __ **EXPOTENTIAL** 04 NARRATIVE DESCRIPTION Mohawk River sediments contaminated by some of substances found at perimeter of site 01 M UNSTABLE CONTAINMENT OF WASTES 02 COSSERVED (DATE. _ POTENTIAL 03 POPULATION POTENTIALLY AFFECTED: 80, 232 04 NARRATIVE DESCRIPTION Former owner allegedly deposited tarry materials on ground, during vehicle cleaning for example. 01 E N DAMAGE TO OFFSITE PROPERTY 02 COBSERVED (DATE. __ ☐ POTENTIAL **04 NARRATIVE DESCRIPTION** None reported 1984 01 & O CONTAMINATION OF SEWERS. STORM DRAINS. WWTPs 02 X OBSERVED (DATE INVESTIGATIONS | POTENTIAL ☐ ALLEGED **04 NARRATIVE DESCRIPTION** Sewers provide preferential pathway for contaminant migration One of most contaminated sites was at outfall of Lee Street sewer in Mohawk 01 P ILLEGAL/UNAUTHORIZED DUMPING 02 C OBSERVED (DATE. _ O POTENTIAL X ALLEGED 04 NARRATIVE DESCRIPTION Tarry wastes allegedly disposed of on or near site 05 DESCRIPTION OF ANY OTHER KNOWN, POTENTIAL, OR ALLEGED HAZARDS None known

☐ ALLEGED

C ALLEGED

☐ ALLEGED

ALLEGED

C ALLEGED

iii. TOTAL POPULATION POTENTIALLY AFFECTED: Approx. 80,232 people within a 3 mile radius IV. COMMENTS

V. SOURCES OF INFORMATION (Cre specific references. • 9 state incs sample analysis, reports)

Harbor Point Property Investigations-Land and River-Niagara Mohawk-1984-1986 Map supplied by Donald S. Youlen, Supervisor, Town of Deerfield to URS Corp. <u>Site Visit</u>

EPA FORM 2070-13 (7-81)

O FRA	POTENTIAL HAZARDOUS WASTE SITE I. IDENTIFICATION						
⊕EPA		SITE INSP	SPECT	TION	<u> </u>	01 STATE 02 SITE NUMBER	
	PART 4 - PERMI	IT AND DES	3CRIP	PTIVE INFORMATI	ION -	NYSDEC #633031	
II. PERMIT INFORMATION	The same of the same	TO BATE I					
01 TYPE OF PERMIT ISSUED (Check at that easy)	02 PERMIT NUMBER	03 DATE IS	SUED	04 EXPIRATION DATE	05 COMMENTS	-	
☐ A. NPDES				 			
□ B. UIC		-			 		
D RCRA	+	+				<u> </u>	
☐ E. RCRA INTERIM STATUS	+						
CXF SPCC PLAN	NA	NA		NA	facility	not in operation	
XCG STATE (Specify) SPDES	0007391	8/1/8		8/1/91	lacino	NOT III OPELACIO	
☐ H LOCAL (Soccity)		 			f		
□ 1. OTHER (Specify)							
□ J. NONE							
III. SITE DESCRIPTION			-				
	2 AMOUNT 03 UNIT 0	OF MEASURE	04 TR	REATMENT (Chock at that as	DOW)	05 OTHER	
☐ A. SURFACE IMPOUNDMENT			I _	INCENERATION		THE NILLIAGE ON GITE	
© B. PILES & C DRUMS, ABOVE GROUND 1111	ıknown			UNDERGROUND INJE		Ø A. BUILDINGS ON SITE	
	known			. CHEMIĈAL/PHYSICAI . BIOLOGICAL			
☐ E. TANK, BELOW GROUND				. BIOLOGICAL . WASTE OIL PROCESS	SMG	06 AREA OF SITE	
☐ F. LANDFELL			□ F. S	SOLVENT RECOVERY	Y	2.96	
G. LANDFARM				OTHER RECYCLING/F	RECOVERY	2.96 (Acres)	
ELOTHER spills un	nknown		D m.	(Spec	icily)	1	
07 COMMENTS				NA			
	-				-		
IV. CONTAINMENT 01 CONTAINMENT OF WASTES (Creek cost)							
	☐ B. MODERATE	🛭 C. INA	ADEQU	JATE, POOR	D. INSECU	RE. UNSOUND, DANGÉROUS	
02 DESCRIPTION OF DRUMS, DIKING, LINERS, BAR	ARIERS, ETC.						
Storage tanks on site.	Allegedly en	mpty. N	lone	known to be	a leaking	. Spills likely	
V. ACCESSIBILITY							
01 WASTE EASILY ACCESSIBLE. AT YES 02 COMMENTS	E NO						
Wastes detected have ma			_	ound.			
VI. SOURCES OF INFORMATION (Che specific							
William Fowlston to Dar Site Visit	1 Rothman, Sep	ot. 16,	1987	7			
<u></u>							

								
		POTI	ENTIAL HAZA			TE		ENTIFICATION
SEPA			SITE INSPEC	TION RE	PORT	,	NY	TATE 02 SITE NUMBER
		PART 5 - WATER	i, Demograph	IIC, AND E	NVIRONN	IENTAL DATA		DEC #633031
II. DRINKING WATER	SUPPLY			····			NYS	DEC HODDOZI
01 TYPE OF DRINKING SUP	PLY		02 STATUS	 				1 DISTANCE TO COM
(Check as applicable)							۱ ~	3 DISTANCE TO SITE
COMMUNITY	SURFACE A.XI	WELL	ENDANGER		ECTED	MONITORED		\
NON-COMMUNITY	C. 🗆	8. 🗆 D. Œ	A. 🗆 D. 25). -	C. 🗆	^	(mi)
	0. [1	0.6	0.10			FO	8	(mi)
III. GROUNDWATER 01 GROUNDWATER USE IN 1	ACDITY (Co				 			
	•							
TA. ONLY SOURCE FO	H ORINKING	© B. DRINKING (Other sources availage COMMERCIAL, IN (No other water source	DUSTRIAL IRRIGATIC		COMMERCIAL Limited other sour	, INDUSTRIAL, IRRIGAT	TON	CI D. NOT USED, UNUSEABLE
02 POPULATION SERVED BY	GROUND WAT	ER 456		03 DISTAN	CE TO NEARE!	ST DRINKING WATER I	WELL	2.8 (mi)
04 DEPTH TO GROUNDWATE	EA .	05 DIRECTION OF GRO	UNDWATER FLOW	OB DEPTH T		07 POTENTIAL YIEL	۵	08 SOLE SOURCE AQUIFER
10-15_	400	North		0FCON		OF AQUIFER		_
				10-1	(8)	Unknown	_ (gpd)	CXYES DINO
09 DESCRIPTION OF WELLS	(including useage,	depth, and location relative to p	population and buildings)	,				
29 Monitori	ng wells	s were insta	lled through	ghout s	urround	ding area f	or n	earby site
						_		eep zones of
		proximately						.00, 2002
10 RECHARGE AREA				11 DISCHAF	~	01		
TYES COMMENTS				XČ YES	COMMENT	•	_	roundwater
□ NO				□ NO	1	discharg	es t	o Mohawk River
IV. SURFACE WATER								
01 SURFACE WATER USE (CA	ect one)							
T 4 55050 4040 000								
A. RESERVOIR, REC	RSOURCE		I, ECONOMICALLY TRESOURCES	Ü C. (COMMERCIA	L. INDUSTRIAL). NOT CURRENTLY USED
							_	
02 AFFECTED/POTENTIALLY	AFFECTED BO	DIES OF WATER						
NAME:						AFFECTED		DISTANCE TO SITE
Mohawk Rive	r					ATEUIED		MSIANCE TO SITE
TIOTICWA REVC.	-							0.2 (mi)
	·				-		_	(mi)
							_	(mi)
V. DEMOGRAPHIC AND	PROPERTY	INFORMATION						
01 TOTAL POPULATION WITH	NN .	7			021	DISTANCE TO NEARES	T PÖPLI	ATION
ONE (1) MILE OF SITE	TW	0 (2) MILES OF SITE	TUDEE /3) MILES OF	1			
A 18.908		19.604	c. 80	, 232	*'E	• C	5	4 4
NO OF PERSONS		NO OF PERSONS		OF PERSONS	-			(mi)
03 NUMBER OF BUILDINGS W	ITHIN TWO (2) A	ALES OF SITE		04 DISTANCE	TO NEAREST	OFF-SITE BUILDING		
•	13.341					.05		
							(m	i)
05 POPULATION WITHIN VICINITY OF SITE (Provide name) of accure of population within vicinity of site, e.g., rurs, village, densely populated under area.								
SITE IIES OF	Site lies on northern side of City of Utica. Largely urban population around site,							
more densely populated to south. Site is on Erie Barge Canal, and is bordered on one side by Utica Harbor.								
one side by	UTICA P	arbor.				•		

POTENTIAL HAZARDOUS WASTE SITE I. IDENTIFICATION **SEPA** SITE INSPECTION REPORT 01 STATE 02 SITE NUMBER PART 5 - WATER, DEMOGRAPHIC, AND ENVIRONMENTAL DATA #633031 VI. ENVIRONMENTAL INFORMATION O1 PERMEABILITY OF UNSATURATED ZONE (CARCIL GRO) □ A. 10⁻⁶ - 10⁻⁶ cm/sec □ B. 10⁻⁴ - 10⁻⁶ cm/sec □ D GREATER THAN 10⁻³ cm/sec 02 PERMEABILITY OF BEDROCK (Creck one) A IMPERMEABLE (Less than 10⁻⁶ cm/sec) © B. RELATIVELY IMPERMEABLE ☐ C. RELATIVELY PERMEABLE ☐ D. VERY PERMEABLE (10⁻⁴ - 10⁻⁶ convenc) (Greater than 10⁻² convenc) 03 DEPTH TO BEDROCK 04 DEPTH OF CONTAMINATED SOIL ZONE 05 SOIL pH unknown 40-85 _(11) unknown 06 NET PRECIPITATION 07 ONE YEAR 24 HOUR RAINFALL OB SLO SITE SLOPE DIRECTION OF SITE SLOPE 14.2 TERRAIN AVERAGE SLOPE 2.3 (in) (in) Variable north-09 FLOOD POTENTIAL 10 east & north-west ☐ SITE IS ON BARRIER ISLAND, COASTAL HIGH HAZARD AREA, RIVERINE FLOODWAY YEAR FLOODPLAIN 11 DISTANCE TO WETLANDS (5 acro months) 12 DISTANCE TO CRITICAL HABITAT (of endangered specific) ESTUARINE ÖTHER 23 ENDANGERED SPECIES none within 1-mile 13 LAND USE IN VICINITY DISTANCE TO: RESIDENTIAL AREAS, NATIONAL/STATE PARKS, FORESTS. OR WILDLIFE RESERVES AGRICULTURAL LANDS COMMERCIAL/INDUSTRIAL PRIME AG LAND AG LAND

14 DESCRIPTION OF SITE IN RELATION TO SURROUNDING TOPOGRAPHY

A. .05

Area has relatively low relief. Urban development (City of Utica) to south. Active and inactive industrial sites in vicinity of site.

.36

none within

none within

VII. SOURCES OF INFORMATION (Can appetite references, e.g., essentia, astropia analysis, reports)

USGS Quadrangle 7.5 minute series, Utica East, N.Y. Leonard E. Ollivett to Linda J. Clark, Aug. 18, 1987 Harbor Point Property Land Investigations-Step 3-March & May 1985

			POTENTIAL HAZARDOUS WASTE SITE	I. IDENTIF	I. IDENTIFICATION		
\$EPA		•	SITE INSPECTION REPORT	OI STATE OF	SITE NUMBER		
		P	NY	11.000.000			
II. SAMPLÉS TAKE	M	<u> </u>	ART 6 - SAMPLE AND FIELD INFORMATION	NYSDEC	#633031		
 		01 NUMBER OF SAMPLES TAKEN	02 SAMPLES SENT TO		03 ESTIMATED DATE		
SAMPLE TYPE		SAMPLES TAKEN			RESULTS AVAILABLE		
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			None taken by Phase I Investiga	tors			
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IV. PHOTOGRAPH							
01 TYPE GROUN	D X AERIAL	(photo copies)	02 IN CUSTODY OF URS Corporation				
03 MAPS	04 LOCATION	OF MAPS		 			
© YES □ NO	URS C	orporation -	570 Delaware Avenue-Buffalo, N.Y.	14202			
V. OTHER FIELD DATA COLLECTED (Provide norrative description)							
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VI. SOURCES OF IN	FORMATIO	N (Cite specific references, e	g state files, sample analysis, regord)				
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O EDA		POTENTIAL HA	NTIAL HAZARDOUS WASTE SITE IL IDENTIFICATION		
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I. CURRENT OWNER(S)		•	PARENT COMPANY (F applicable)	- NYSDE	C #633031
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III. OFF-SITE GENERATOR(S)	<u>t</u>	<u> </u>					
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BEPA	POTENTIAL HAZARDOUS WASTES SITE INSPECTION REPORT		L IDENTIFICATION
	PART 10 - PAST RESPONSE ACTIVITI	ES ,	NY YSDEC #633031
PAST RESPONSE ACTIVITIES			12DEC #633031
01 A. WATER SUPPLY CLOSED	02 DATE	03 AGENCY	
04 DESCRIPTION			
01 [] B. TEMPORARY WATER SUPPLY PR 04 DESCRIPTION	ROVIDED 02 DATE	03 AGENCY	
01 C. PERMANENT WATER SUPPLY PR 04 DESCRIPTION	POVIDED 02 DATE	03 AGENCY _	
01 [] D. SPILLED MATERIAL REMOVED 04 DESCRIPTION	02 DATE	03 AGENCY	
01 G E. CONTAMINATED SOIL REMOVED 04 DESCRIPTION	OZ DATE	03 AGENCY _	
01 [] F. WASTE REPACKAGED 04 DESCRIPTION	OZ DATE	03 AGENCY _	
01 G. WASTE DISPOSED ELSEWHERE	O2 DATE	O3 AGENTY	
04 DESCRIPTION		************************************	
01 D H. ON SITE BURIAL 04 DESCRIPTION	02 DATE	O3 AGENCY _	
01 (I. IN SITU CHEMICAL TREATMENT 04 DESCRIPTION	O2 DATE	03 AGENCY _	
01 [] J. IN SITU BIOLOGICAL TREATMENT 04 DESCRIPTION	O2 DATE	03 AGENCY _	
01 [] K. IN SITU PHYSICAL TREATMENT 04 DESCRIPTION	02 DATE	03 AGENCY	
01 C L ENCAPSULATION 04 DESCRIPTION	02 DATE	03 AGENCY	
01 D M EMERGENCY WASTE TREATMENT	O2 DATE	03 AGENCY	
04 DESCRIPTION			
01 D. CUTOFF WALLS 04 DESCRIPTION	02 DATE	03 AGENCY	
01 C O. EMERGENCY DIKING/SURFACE WAS 04 DESCRIPTION	ATER DIVERSION 02 DATE	_ 03 AGENCY	
01 DP CUTOFF TRENCHES/SUMP 04 DESCRIPTION	02 DATE	03 AGENCY	
01 C Q. SUBSURFACE CUTOFF WALL	02 DATE	03 AGRIGO	

OFDA	POTENTIAL HAZARDOUS WASTE SITE	İ	I. IDENTIFICATION OF STATE OF SITE NUMBER
\$EPA	SITE INSPECTION REPORT PART 10 - PAST RESPONSE ACTIVITIES		NY
II PAST RESPONSE ACTIVITIES (Comman)	PANT IV-PAST RESPONDE AUTIVITIES		NYSDEC #633031
01 C R BARRIER WALLS CONSTRUCTED	2000		
04 DESCRIPTION	02 DATE		
01 🗆 S. CAPPING/COVERING 04 DESCRIPTION	02 DATE		
01 🗍 T. BULK TANKAGE REPAIRED 04 DESCRIPTION	O2 DATE		
01 U. GROUT CURTAIN CONSTRUCTED 04 DESCRIPTION	02 DATE		-
01 🗆 V. BOTTOM SEALED 04 DESCRIPTION	02 DATE		
01 C W GAS CONTROL 04 DESCRIPTION	02 DATE		
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01 [] 2. POPULATION RELOCATED 04 DESCRIPTION	O2 DATE	03 AGENCY_	,
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III. SOURCES OF INFORMATION (Che specific return	Micros. 4.9 . Mileto (Res. semple enalysis, reports)		
NYSDEC - Region 6			
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POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 11 - ENFORCEMENT INFORMATION

L IDENTIFICATION

01 STATE 02 SITE NUMBER

NY

NYSDEC #633031

II. ENFORCEMENT INFORMATION

01 PAST REGULATORY/ENFORCEMENT ACTION | YES | XI NO

02 DESCRIPTION OF FEDERAL, STATE, LOCAL REGULATORY, ENFORCEMENT ACTION

III. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sarrole energies, reports)

NYSDEC Region 6

EPA FORM 2070-13 (7-81)

6.0 ASSESSMENT OF DATA ADEQUACY AND RECOMMENDATIONS

Data collected during this Phase I investigation of the New York Emulsion Tar Products site are considered inadequate in the following areas:

o Waste characteristics:

- Data should be developed and evaluated concerning the derivation of tars and oils found in soils on and adjacent to the site -- i.e., data indicating whether the tars and oils are coal-derived or petroleum-based.
- Soil borings and monitoring wells should be installed on the site itself, with sampling and analysis of soil and groundwater.

6.1 APPENDIX A - DATA SOURCES AND REFERENCES

REFERÊNCES - DOCUMENTATION RECORDS

- 1. Harbor Point Property Land Investigations, Results of Extended Site Investigations, Step 3 Land Report, Niagara Mohawk Power Corporation, March 1985 (Revised May 1985).
- 2. Harbor Point Property River Investigations, Remedial Design Concepts, Step 4 River Report, Niagara Mohawk Power Corporation, June 1986.
- 3. Uncontrolled Hazardous Waste Site Ranking System, A Users Manual (HW-10), United States Environmental Protection Agency 1984, Figures 4 and 8, Tables 2, 3, 4, 5, and 9, pp. 24, 25, and 27.
- 4. Dangerous Properties of Industrial Materials Sixth Edition, N. Irving Sax, Van Nostrand Reinhold Company, New York, 1984.
- 5. Máp and Statement by Donáld S. Youlen, Supérvisor, Town of Deerfield, New York, October 26, 1987.
- 6. Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, to Russell Logalbo, Utica Board of Water Supply, September 10, 1987.
- 6a. Karl P. Maxwell, Supervisor, Town of Marcy, New York, to Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, November 12, 1987.
- 7. Robin Mangini, District Conservationist, United States Department of Agriculture, Soil Conservation Service, to Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, September 14, 1987.
- 8. USGS Topographic Maps, 7.5 Series: Utica East, New York, Quadrangle, 1983; Utica West, New York, Quadrangle, 1955; South Trenton, New York, Quadrangle, 1983; Oriskany, New York, Quadrangle, 1955.

- 9. City of Utica, Oneida County, New York Tax Map, Sheet No. 318.08.
- 10. City of Utica, Oneida County, New York Aerial Maps, Sheet Nos. 306, 318.
- 11. William Fowlston, Safety Director, Suit-Kote Corporation, to Daniel W. Rothman, Phase I Investigation Project Manager, URS Corporation, September 16, 1987.
- 12. Harbor Point Property Land Investigations, Proposal for Initial Site Survey, Step 1 Land Report, Niagara Mohawk Power Corporation, March 1984.
- 13. Muffett A. Mauche, Staff Engineer, LeRoy Callender, PC, telecon to James Doyle, Sanitary Engineer, NYSDEC Region 6, October 8, 1987.
- 14. Leonard E. Ollivett, Conservation Biologist, NYSDEC Region 6, to Linda J. Clark, Project Geologist, URS Corporation, August 18, 1987.
- 15. New York State Atlas of Community Water System Sources, New York State Department of Health, Division of Environmental Protection, Bureau of Public Water Supply Protection, 1982.
- 16. Telecon Karen A. Hartnett, URS Corporation, to Fire Chief, City of Utica, September 2, 1987.
- 17. 1980 Census of Population, Number of Inhabitants, New York, United States Department of Commerce, Bureau of the Census.
- 18. Harbor Point Property: The Mohawk River, Proposal for Initial River Survey, Step 1 River Report, Niagara Mohawk Power Corporation, March 1984.

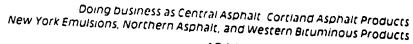
- 19. William Fowlston, Safety Director, Suit-Kote Corporation, to Kenneth J. Crandall, Jr., Chief, Spill Prevention Section, Division of Water, NYSDEC, June 22, 1987.
- 20. Harbor Point Property Land Investigations, Results of Initial Survey, Step 2 Land Report, Niagara Mohawk Power Corporation, August 1984.
- 21. Geology of New York: A Short Account, New York State Museum and Science Service, Educational Leaflet No. 20, 1966.
- 22. Geologic Map of New York, Hudson-Mohawk Sheet, 1970.
- 23. New York State Geological Association Guidebook, 36th Annual Meeting, May 8-10, 1964, Department of Geology, Syracuse University.
- 24. Stuart E. Smith, Sr. Sanitary Engineer, NYSDEC Region 6, to James Luz, Regional Water Engineer, NYSDEC Region 6, March 11, 1985, with attachments.
- 25. Harbor Point Property: The Mohawk River, Results of Initial River Survey, Step 2 River Report, Niagara Mohawk Power Corporation, September 1984.
- 26. Miscellaneous References on Nature of Chemical Materials and Processes, various sources.
- 27. Climates of the States, New York, Climatography of the United States, No. 60-30, United States Department of Commerce, Weather Bureau, February 1960.

- 28. Inactive Hazardous Waste Disposal Site Report, with attachments, November 21, 1986.
- 29. Darrell Sweredoski, Sr. Sanitary Engineer, NYSDEC Region 6, to Walt Demick, July 14, 1987.
- 30. New York State Department of Environmental Conservation Report Inactive Hazardous Waste Disposal Sites in New York State, Volume 6, 1989. Site Reports for: Niagara Mohawk Harbor Point Property (633021), Monarch Chemical Company, Inc. (633030), and Mohawk Valley Oil, Inc. (633032).

EPA 2070-13

HARBOR POINT PROPERTY: THE MOHAWK RIVER
PROPOSAL FOR INITIAL RIVER SURVEY
STÉP 1 ŘÍVÉR REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York





1911 Lorings Crossing Road P.O. Box 5160 Cortland, N.Y. 13045-5160

REF. (19)

Walkins Glen

607 535 2743

June 22, 1987

NYSDEC Spill Prevention Section 50 Wolf Road, Room 300 Albany, NY 12233

Attention: Mr. Kenneth J. Crandall, Jr. P.E. Chief, Spill Prevention Section Division of Water

Dear Mr. Crandall:

In regards to your letter of 6/12/87 to Rod Birdsall, President of Suit-Kote, I wish to explain our current position regarding the location at Utica. We ceased all operations at that plant in the fall of 1984. The following summer we removed all products and dismantled the plant.

On Wednesday, June 17, 1987, I met with a Mr. Mark Kepple of Environmental Oil at the location. The purpose of our inspection was to establish an inventory of tanks and equipment in order for Environmental to present us an estimate for removal and clean up at the yard. I expect to obtain a total of three estimates on the clean up operations before we actually start the project.

I will keep you posted on our progress.

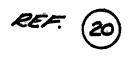
Bill fowlston

Bill Fowlston

Director of Safety and Environment

dr

Buffalo Cortland McGraw 716-683-8850 607-753-6085 New Harrlord Norwich 607 749-2665 607-753-3358 W Oneonta 315-735-8501 Rochester 607-336-7112 Vestal Watertown 607-432-0440 716-473-6321 607 729 1001 315-782 4457



HARBOR POINT PROPERTY LAND INVESTIGATIONS RESULTS OF INITIAL SURVEY STEP 2 LAND REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York

REF. (21)

Geology of New York: a short account

Adapted from the text of "Geologic Map of New York State" by J. G. Broughton, D. W. Fisher, Y. W. Isachsen, and L. V. Rickard

EDUCATIONAL LEAFLET NO. 20

The University of the State of New York/The State Education Department
New York State Museum and Science Service/Albany 1966

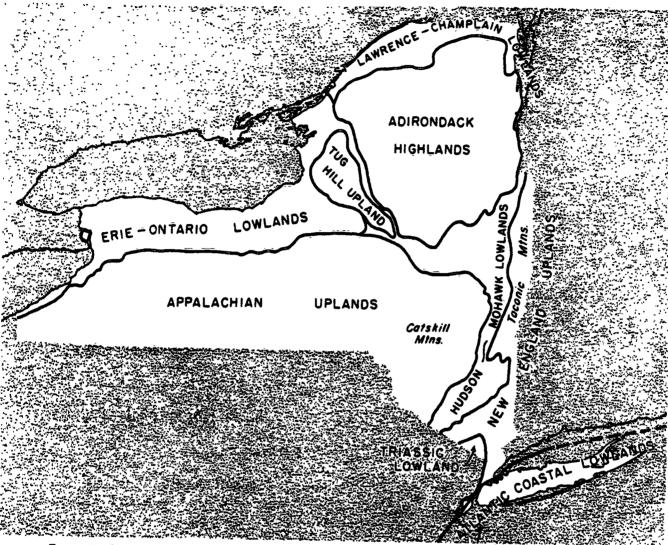


Figure 19. Physiographic provinces of New York, based on relief and geology (Modified after G. B. Cressey, 1952)

Cenozoic Era

PHYSIOGRAPHIC PROVINCES AND TERTIARY HISTORY

The physiographic provinces of New York are shown in figure 19. Modern landscapes of the State were shaped largely during the Cenozoic Era, the most recent 65 million years of geologic history. Although the overall features later would be modified and blurred by glaciation, the broad outlines of modern mountain, valley, and plain first were carved by the unrelenting rush of water to the earlier Cenozoic seas.

The long sequence of erosion presumably began with the arching of the Jurassic Fall Zone erosion surface in mid-Cretaceous time. As its eastern flank dipped beneath the encroaching Atlantic Ocean to receive Coastal Plain deposits, the axis domed sufficiently to initiate the sculpture of the Appalachians and Adirondacks. Few, if any of today's land forms can be traced so far back, however. Most researchers believe that all the exposed remnants of the dissected Fall Zone surface were obliterated by subsequent erosion.

South of New York, at least a partial record of Tertiary geology persists in the Coastal Plain deposits. In addition to a sedimentary record, datable igneous intrusions cut rocks of varying degrees of deformation in the western states. But in New York, no such tangible evidence of Cenozoic events exists. The Coastal Plains sediments derived from the long-continued degradation of New York and New England now rest on the Continental

Shelf, beneath many fathoms of water. Because of a relatively recent tilting of the coastline about a northwest-southeast axis near New York City, the Coastal Plain has been raised south of New York; east and north of the city, all but the Long Island Cretaceous has been depressed below sea level.

Since exposed Tertiary sedimentary deposits are absent in New York, its geological history must be reconstructed from the only data available, the present physiographic features of the State. In an area as small as New York, where climate does not vary significantly, land forms have been determined primarily by geology. Characteristic differences between the physiographic provinces have resulted from the ways in which rocks of differing lithologies and structures have reacted to the erosional force of the Cenozoic. Thus, while many authorities have classified New York's physiographic provinces in various ways, all are more or less in agreement as to the outlines of the major provinces; they differ mainly in the names applied to the provinces. Those used here were proposed by George B. Cressey (1952, personal communication, J.G. B.). From north to south, the physiographic provinces of New York are:

St. Lawrence-Champlain Lowlands

New York's northernmost province includes the St. Lawrence River Valley (northeast of the Thousand Islands), the low hills south of the river valley, and the Lake Champlain Valley (figure 19). The underlying rocks—Cambrian and Ordovician sandstones, dolomites, and limestones—dip gently away from the Adirondacks. Relief is approximately 100 feet. Streams draining the northern and eastern slopes of the Adirondacks flow across the province. The shoreline of Lake Champlain is largely controlled by north-south and east-west faults which have chopped the Paleozoic sandstones and carbonates into large blocks.

Adirondack Highlands

The highest mountains in New York occur in the Adirondack Highlands, especially in the High Peaks region; the High Peaks, in the east-central part of the province, are underlain by anorthosite, which is highly resistant to erosion. Two peaks—Mt. Marcy and Mt. Algonquin—are over 5,000 feet in elevation, and many exceed 4,000 feet. Average relief in the Adirondack Highlands is 2,000 feet. North, west, and south of the High Peaks area, elevations decrease gradually; east to the Champlain Lowland, the slope is more abrupt.

The Adirondacks are transected by long, northeast-southwest lineaments, representing shear zones or major faults. The lineaments frequently control drainage and the shape of land forms. Many lakes follow geologic contacts, or are confined to valleys along weak metasedimentary rocks. Because glacial deposits have clogged the normal radial drainage, lower areas are dotted with lakes, ponds, and swamps.

Tug Hill Upland

The Tug Hill, an isolated upland in the eastern part of the Erie-Ontario Lowlands, is probably the most desolate area of the State. Elevation is 1,800 to 2,000 feet, and relief is very low. The Tug Hill results from a resistant cap rock of Oswego Sandstone (an Ordovician sedimentary quartzite), resting on a thick series of sandy shales. These, in turn, overlie Trenton and Black River limestones, which form a flight of rock terraces along the west side of the Black River Valley. The low slope of the cap rock and the thin cover of glacial deposits have caused poor drainage and many swamps.

Erie-Ontario Lowlands

This province encompasses the relatively low, flat areas lying south of Lake Erie and Lake Ontario and extending up the Black River Valley. From the lake levels of 570 feet and 244 feet, respectively, the land rises gently eastward and southward. The maximum elevation (1,000-1,500 feet) occurs along the Portage Escarpment, the boundary with the Appalachian Uplands to the south. Particularly in the Ontario Lowland, east-west escarpments are formed by the Onondaga Limestone and Lockport Dolomite. (The Lockport is the cap rock of Niagara Falls and the falls of the Genesee River at Rochester.) The simple erosional topography has been modified substantially by glacial deposition of drumlin fields, recessional moraines, and shoreline deposits.

Hudson-Mohawk Lowlands

The general topography of the Hudson-Mohawk Lowlands resulted from erosion along outcrop belts of weak rocks. In the Mohawk Lowlands, the outcrop belts lie between the Adirondacks and the Helderberg Escarpment; for the Hudson, they lie between the Catskills and the metamorphosed shale hills of the Taconics. Most of the province has low elevation and relief. It is underlain primarily by Ordovician shales which have been exposed by the southward and westward stripping off of Silurian and Devonian limestones.



COBLESKILL LIMESTONE AND SALINA GROUP 0-700 ft. (0-210 m.)

Cobleskill Limestone; Bertie, Camillus, and Syracuse Formations—shale, dolostone; Brayman Shale.

Sv Vernon Shale.

Scs

Scl

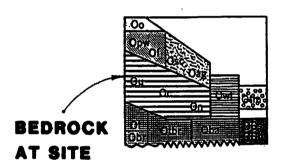
LOCKPORT GROUP 0-75 ft. (0-23 m.)

SI Ilion Shale.



CLINTON GROUP 0-350 ft. (0-110 m.)

Herkimer Sandstone including Joslin Hill and Jordanville Members; Kirkland Hematite; Willowvale Shale; Westmoreland Hematite; Sauquoit Formation—sandstone, shale; Otsquago Sandstone; Oneida Conglomerate.



LORRAINE, TRENTON, AND BLACK RIVER GROUPS up to 4,500 ft. (1400 m.)

Oo Oswego Sandstone

Opw Pulaski and Whetstone Gulf Formations—shale, siltstone.

Of Frankfort Formation—shale, siltstone.

Osc Schenectady Formation—graywacke, sandstone, siltstone, shale.

Oag Austin Glen Formation-graywacke, shale.

Ou Utica Shale.

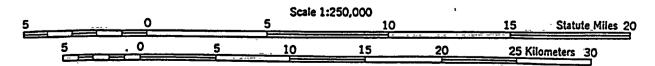
Oc Canajoharie Shale.

On Normanskill Shale—minor mudstone, sandstone.

GEOLOGIC MAP OF NEW YORK

1970

Hudson-Mohawk Sheet



CONTOUR INTERVAL 100 FEET

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145 N56 1964

NEW YORK STATE GEOLOGICAL ASSOCIATION

36th ANNUAL MEETING MAY 8-10, 1964

GUIDEBOOK



DEPARTMENT OF GEOLOGY, SYRACUSE UNIVERSITY



by Irwin H. Kantrowitz

U.S. Geological Survey

All but the largest public water systems in the Syracuse area obtain their supply from wells or springs. Almost all farms and homes in rural areas are supplied by private wells or springs and many industries also rely on ground-water supplies to meet their needs. Current withdrawal of ground water in the area is believed to be only a fraction of the available supply. The quality of water, however, is not always suitable for many uses, including public supply.

Ground water occurs in fractures and bedding joints of consolidated rocks and in pore spaces of unconsolidated deposits. The quantity of water available depends on the nature of the aquifer and the source of recharge. Adequate supplies for domestic and farm needs (100 to 1,000 gallons per day) are almost always available. Larger quantities of water for industrial and public supplies can generally be obtained from stratified coarse-grained deposits and, less frequently, from bedrock with prominent fractures, particularly where these aquifers are in hydraulic contact with a surface-water body which acts as a source of recharge. Ground-water quality depends on the chemical characteristics of the aquifer material, and flow pattern within the ground-water reservoir, and the quality of the recharge water. The factors most commonly affecting the quality of the ground water in the Syracuse area are hardness, iron, hydrogen sulfide, and salinity.

Ground water in consolidated rocks

Table I shows the rock units in the Syracuse area, their dominant lithologies, and the quality of the ground water that may be expected in wells tapping each unit. Wells in the limestone units, and the Camillus Shale, Syracuse Salt, and Vernon Shale will yield as much as 230 gpm (gallons per minute) because of enlargement of fractures by the solution of the carbonates and evaporites. The yield of wells drilled in these units for domestic, farm, and other small supplies averages about 15 to 20 gpm. Wells in the other rock units in the area generally yield less than 10 gpm and are inadequate for most public or industrial needs.

Carbonate (temporary) hardness results from the solution of limestone or dolomite by ground water. The hardness of water in the Camillus and Vernon Shales is predominantly noncarbonate (permanent) hardness resulting from the solution of gypsum or anhydrite. The source of hydrogen sulfide is believed to be pyrite found in the Hamilton Group and the Lorraine and Utica Shales, and sphalerite found in the Lockport Dolomite. Although traces of iron are found in water from all the rock units, it is present in objectionable concentrations most often in the Camillus and Vernon Shales where it is probably related to the occurrence of hematite, siderite, and pyrite.

The presence of saline water (here defined as water containing more than 250 parts per million of chloride) is not shown in Table 1 because its occurrence is more closely related to patterns of ground-water movement than it is to

Data contained in this summary were collected by the U.S. Geological Survey in cooperation with the New York State Water Resources Commission. Publication authorized by Director, U.S. Geological Survey.

the chemical characteristics of the water-bearing units. Although the only salt beds in the area are found within the Syracuse, most wells tapping this formation in its outcrop area do not yield salty water because the salt at shallow depths has been almost completely dissolved. Wells drilled into the Syracuse in the area south of its outcrop generally yield saline water, and commercial brine is obtained from deep wells in Tully Valley, about 12 miles south of Syracuse. At these wells, the salt occurs 300 to 500 feet below sea level and the brine is produced by injecting fresh water into the beds and then pumping it out after it has dissolved the salt.

The major area of natural saline-water occurrence is along the lowlands occupied by Oneida Lake and the Oneida, Oswego, and Seneca Rivers. This area coincides with the major area of ground-water discharge and the presence of saline water is believed to be due to the upward and northward movement of ground water that has been in contact with and partially dissolved the salt beds beneath the Appalachian Plateau. Wells drilled more than 100 feet into the Genesee Formation or Hamilton Group in the valleys of the plateau area may also yield saline water. The occurrence of this water may be related to connate water within the rock units or to the upward movement of water from the salt beds.

Ground water in unconsolidated deposits

A till sheet commonly about 30 feet thick mantles the entire upland area in the Appalachian and Tug Hill Plateaus and a large part of the Ontario lowland. Adequate supplies of water for domestic and farm supplies are generally available from dug wells or springs, although shallow wells on hillsides and hilltops frequently are inadequate during long dry periods.

Stratified drift mantles the remainder of the area, notably in the valleys of the Appalachian Plateau, most of the Ontario Lowland, and the lower parts of the valleys of the Tug Hill Plateau. Deposition of stratified drift occurred under four conditions: 1) proglacial deposition during free drainage, 2) deposition in ice-dammed valleys, 3) deposition during Great Lakes drainage, and 4) deposition in Lake Iroquois.

Coarse-grained glaciofluvial deposits consisting largely of sand and gravel occur south of the Valley Heads moraine and in many places form a large part of the moraine itself. The sand and gravel are well sorted and are probably the most permeable water-bearing material in the area. The city of Cortland, located about 27 miles south of Syracuse and 14 miles south of the Valley Heads moraine, pumps more than 2.5 mgd (million gallons per day) from these deposits. Somewhat similar sands and gravels, deposited during free glacial drainage in the Tug Hill Plateau area, may be expected along West Branch Fish Creek.

During deglaciation of the Appalachian Plateau, lakes existed in the major valleys, dammed between the bedrock divide to the south and the ice tongue to the north. Although data are scanty, the deposits in the valleys appear to become coarser with increasing depth which is consistent with a concept of a receding source of sediment. Small but adequate domestic and farm supplies can generally be obtained from wells dug in lacustrine sand, silt or clay, and driven screened wells are common where lacustrine sands occur at shallow depths. Because the layers of gravel in these deposits are lenticular, few wells drawing from gravelyield more than 100 gpm and the average yield of such wells is only about 30 gpm.

With further deglaciation, the ice margin was against the escarpment of the

(23)

Appalachian Plateau, and eastward drainage of the ancestral Great Lakes was initiated in ice-marginal channels. Deposition of sand and gravel occurred whereever the Great Lakes waters entered standing water in the north-south valleys or where westward recession of the ice front enabled the water to abandon the marginal channels and utilize the larger north-south valleys as outlets to the lowland north of the escarpment. These sand and gravel deposits are probably not as permeable as the valley train material south of the Valley Heads moraine. They are, nevertheless, a potential source of large ground-water supplies because they generally occur in areas where stream infiltration is possible. Examples of wells in this type of deposit are a public-supply well for the village of Fayetteville that has been test pumped at 500 gpm, and a public-supply well for the village of Chittenango that yields 350 gpm.

During the last stages of deglaciation in the Syracuse area, Lake Iroquois, a proglacial ancestral Lake Ontario, occupied the lowland north of the Appalachian escarpment. Melt-water streams deposited outwash deltas in the lake which were subsequently reworked and covered by finer grained lacustrine deposits as the ice continued to recede. The sand and gravel, where it is in hydraulic contact with a surface-water body may yield large quantities of water. The village of Fulton has pumped as much as 3.3 mgd from a well field adjacent to the Oswego River. Individual wells in this system yield as much as 800 gpm.

For the most part, none of the unconsolidated deposits in the Syracuse area have undergone significant transport by ice or melt water. Therefore, the chemical nature of the deposits and, to a large measure, the quality of the ground water derived from them, is generally similar to that of the underlying bedrock. Saline water occurs notably in a few of the north-south valleys where ground water has been able to move from the truncated salt beds of the Syracuse into relatively permeable valley-fill material.

Table i.--Water-bearing units and quality of ground water

Rock unit	Lithologic type	Quality of water
Genesee Formation	shale	generally good
Tully Limestone	limestone	hard
Hamilton Group	shale, limestone	hard, hydrogen sulfide
Onondaga Limestone	1 imes tone	hard
Helderberg Group	limestone	hard
Cobleskill Limestone	l imes tone	hard
Bertie Limestone (of Salina Group)	limestone, dolomite, some shale	hard
Camillus Shale (of Salina Group)	shale, gypsum, dolomite	hard, iron
Syracuse Salt (of Salina Group)	shale, gypsum, dolomite, salt	hard, iron
/ernon Shale (of Salina Group)	shale, some gypsum & dolomite	hard, iron
Lockport Dolomite	dolamite	hard, hydrogen sulfide
Clinton Group	sandstone & shale, some limestone	hard
Albion Group	sandstone	generally good
Queenston Shale	sands tone	generally good
swego Sandstone	sands tone	generally good
eorraine Shale	shale	hydrogen sulfide
Utica Shale	shale	hydrogen culfide
Approximately equivalent to	Medina Group of N.Y. State	Geological Survey usage





New York State Department of Environmental Conservation

MEMORANDUM

TO:

Jim Luz

FROM:

Stuart E. Smith

SUBJECT:

Sample Results R-684-045-01 & 02, New York Emulsions (Koppers),

•

Utica (C), Oneida County

DATE: March 11, 1985

On 29 November 1984, I took a sample (-01) of tar from the ground beside a tar tank on the subject property. The tank was on the NE side of the property, near the dyke, near the chain-link fence line which is on the Emulsions/Niagara Mohawk property line. A second sample (-02) was taken on the Niagara Mohawk property some 5-10 feet from the fence.

The analytical results are attached. The most significant parameters present in the samples were

naphthalene
2-methylnaphthalene

at about the same concentration in each sample.

The results suggest that the sample collected on Niagara Mohawk property may have originated from the New York Emulsions property, provided that property ownership has not been changed over the last 10 to 15 years. The tar tank (sample -01) was last used about 10 years ago, according to Jim Cromie, a New York Emulsions employee.

Stuart E. Smith, P. E. Sr. Sanitary Engineer

Region 6 - Utica

Att. SES:m

cc: John Kenna, w/attachments

GCA/TECHNOLOGY DIVISION A DIVISION OF GCA CORPORATION Bedford, Massachusetts 01730

CERTIFICATION

STATE OF MASSACHUSETTS)
COUNTY OF MIDDLESEX)ss:

I, Deborah McGrath, Analytical Chemist and Operations Manager of the Laboratory Analysis Department of GCA/Technology Division, an Analytical Laboratory, do hereby certify that the annexed item is a true and correct copy of an original Analytical Laboratory report, Identification No. GCA 5-668-073, the transmission letter of which is dated 19 February 1985.

The original of this report is on file in the Analytical Laboratory located at 213 Burlington Road, Bedford, MA 01730.

Deborah McGrath(Analytical Chemist Operations Manager Laboratory Analysis Dept.

Sworn ic before re this 19th day of February 1985.

Paul G. Merrill Notary Walic

My Commission Engines: 12/27/66

	Contractor	GCA/Technology	y Division
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NYDEC Contract'# __C000678. .

Narrative For GCA 5-668-073

SAMPLE RECEIPT

- 1. Two (2) samples were received by GCA/Technology Division on December 18, 1984. The requested analyses for the individual samples are provided in Table 1.
- 2. Upon receipt, the submitted samples were entered in the Master Log Book and assigned GCA Control Numbers as presented in Table 1.

ANALYTICAL PROCEDURES

1. A 0.5 gram aliquot of each sample was diluted in methylene chloride for GC/MS analysis. Both sample aliquots required dilution with 1000 ml of solvent in order to provide a free-flowing solution suitable for injection. Analysis was performed using GC/MS techniques as specified in EPA Method 625. The results of analyses are provided on the attached Data Report Sheets. Also provided are copies of the reconstructed ion chromatograms (RICs) which have been labeled for internal standards.

QUALITY CONTROL

 Quality control samples appropriate for this analysis are not available, therefore none were analyzed. Surrogate spiking was not performed because the samples required a large diluent volume.



TABLE 1. CROSS REFERENCE LIST OF SUBMITTED SAMPLES

GCA Control No.	Sample . Receipt	Sample Matrix	Sample Identification	Requested . Analyses
42239	12/18/84	Tar	R-684-045-01	Base/Neutral and Acid Extractable Organics
42240	•		R-684-045-02	



Project <u>5-668</u> -	-073
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Sample I.D. R-684-045-01

GCA Control No. 42239

1/29/85

DATA REPORT SHEET

Base/Neutral Extractables (Page 1 of 2)

Sample I.D. $R-684-045-01$	Analysis Date	1/29/85	
Sample Matrix Tar	Instrument HP 5985 GC/MS		
Parameter	Ion Used To Quantitate	Concentration (mg/kg)	
N-nitrosodimethylamine		ND	
bis(chloromethyl)ether		ND ND	
bis(2-chloroethyl)ether		ND ND	
1.3-dichlorobenzene		ND	
1,4-dichlorobenzene	_	ND ND	
1,2-dichlorobenzene		ND	
bis(2-chloroisopropyl)ether		ND ND	
N-nitrosodi-n-propylamine		ND .	
hexachloroethane		ND ND	
nitrobenzene		ND ND	
isophorone		ND	
bis(2-chloroethoxy)methane		ND	
1,2,4-trichlorobenzene		ND	
naphthalene	128	12000	
hexachlorobutadiene		ND	
hexachlorocyclopentadiene	,	ND	
2-chloronaphthalene		ND	
dimethyl phthalate		ND ND	
acenaphthylene		ND ND	
2,6-dinitrotoluene	_	ND	
acenaphthene		ND	
2,4-dinitrotoluene		ND**	
diethyl phthalate		ND ND	
fluorene		ND*	
4-chlorophenyl phenyl ether		ND	
N-nitrosodiphenylamine		ND	
1,2-diphenylhydrazine		ND ND	
4-bromophenyl phenyl ether		ND*	
hexachlorobenzene		ND*	
phenanthrene/anthracene d		ND	
di-n-butyl phthalate		175	

ND = <10,000 mg/kg ND* = <40,000 mg/kgND** = < 100,000 mg/kg





Project	5-668-073	001 0 6 /0000	
		GCA Control No. 42239	

DATA REPORT SHEET

Base/Neutral Extractables (Page 2 of 2)

Sample I.D. R-684-045-01	Analysis Date	1/29/85
Sample Matrix Tar Instrume	nt <u>HP 5985 (</u>	GC/MS
Parameter	Ion Used To Quantitate	Concentration (mg/kg)
fluoranthene		ND*
benzidine		ND**
pyrene		ND
butyl benzyl phthalate		ND*
3,3'-dichlorobenzidine		ND*
benzo(a)anthracene/chrysene ^a		ND .
bis(2-ethylhexyl)phthalate		ND
di-n-octyl phthalate		ND*
benzo(b)fluoranthene/benzo(k)fluoranthene		ND*
benzo(a)pyrene		ND*
indeno(1,2,3-cd)pyrene		ND*
dibenzo(a,h)anthracene		ND*
benzo(g,h,i)perylene		ND*
2,3,7,8-tetrachlorodibenzo-p-dioxin		ND**

 $\begin{array}{ll} ND = < 10,000 \text{ mg/kg} \\ ND* = < 40,000 \text{ mg/kg} \\ ND** = < 100,000 \text{ mg/kg} \end{array}$

^aUnder the conditions stipulated by EPA Method 625 this compound pair cannot be chromatographically resolved. The concentration reported reflects the total of both isomers.





GCA Control No. 42239

DATA RÉPORT SHEET

Acid Extractables

Analysis Date 1/29/85			
Instrument	HP 5985 GC/MS		
Ion Used To Quantitate	Concentration (mg/kg)		
•	ND		
	ND**		
	Instrument Ion Used ToQuantitate		

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ND = < 10,000 mg/kgND** = < 100,000 mg/kg



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Project_	5-668-073

GCA	Control	No.	42239
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DATA REPORT SHEET

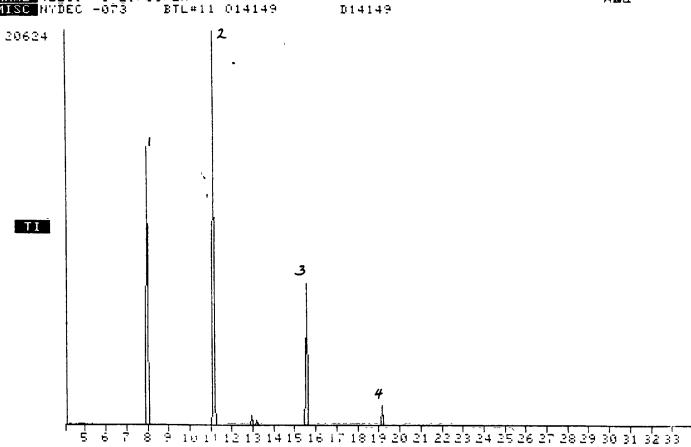
Extractables Qualitative Analysis

Sample I.D. R-684-045-01		Analysis Date 1/29/85		
Sample Matrix Tar	Instrument_	HP 5985 GC/MS		
Compound		CAS No.	Concentration* (mg/kg)	
2-methylnaphthalene	-	91-57-6	10,000	
		,		

^{*}Concentration based on a comparison of the total ion area of the compound with that of the internal standard.

मृत्रश्रम् ।

NAME 42239 1 29.85.EM MISC NYDEC -073 BTL#11 014149



- 1 = D₄-1,4-dichlorobenzene
- 2 = Dg-naphthalene
- $3 = D_{10}$ -acenaphthene
- $4 = 0_{10}$ -phenanthrene

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Project	5-668-073	
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GCA Control No. 42240

DATA REPORT SHEET

Base/Neutral Extractables (Page 1 of 2)

Sample I.D. R-684-045-02	Analysis Date _	1/29/85
Sample Matrix Tar	Instrument HP 5985 G	C/MS
Parameter	Ion Used To Quantitate	Concentration (mg/kg)
N-nitrosodimethylamine		ND ND
bis(chloromethyl)ether		ND ND
bis(2-chloroethyl)ether		ND
1.3-dichlorobenzene		ND ND
1.4-dichlorobenzene		ND
1,2-dichlorobenzene		ND.
bis(2-chloroisopropyl)ether		- ND
N-nitrosodi-n-propylamine		ND .
hexachloroethane		- ND
nitrobenzene		ND
isophorone		ND
bis(2-chloroethoxy)methane		ND
1,2,4-trichlorobenzene		ND
naphthalene	128	19000
hexachlorobutadiene		ND _
hexachlorocyclopentadiene		ND
2-chloronaphthalene		ND
dimethyl phthalate		ND
acenaphthylene		ND
2,6-dinitrotoluene		ND
acenaphthene		ND
2,4-dinitrotoluene		ND**
diethyl phthalate fluorene		ND.
		ND
4-chlorophenyl phenyl ether		ND*
N-nitrosodiphenylamine		ND
1,2-diphenylhydrazine		ND
4-bromophenyl phenyl ether hexachlorobenzene		ND*
		ND*
phenanthrene/anthracene a		ND
di-n-butyl phthalate		ND

ND = < 10,000 mg/kg ND* = < 40,000 mg/kg ND** = < 100,000 mg/kg





Project	5-668-073
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GCA	Control	No.	42240
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DATA REPORT SHEET

Base/Neutral Extractables (Page 2 of 2)

Sample I.D. R-684-045-02	Analysis Date _	1/29/85
Sample Matrix Tar Instrume	ntHP 5985 (GC/MS
Parameter	Ion Used To Quantitate	Concentration (mg/kg)
fluoranthene		ND*
benzidine		
pyrene		ND**
butyl benzyl phthalate		ND
3,3'-dichlorobenzidine		ND*
benzo(a)anthracene/chrysene ²		ND*
bis(2-ethylhexyl)phthalate		ND .
distribution of the state of th		ND .
di-n-octyl phthalate		ND*
benzo(b)fluoranthene/benzo(k)fluoranthenea		ND*
benzo(a)pyrene		ND*
indeno(1,2,3-cd)pyrene		
dibenzo(a,h)anthracene		ND*
benzo(g,h,i)perylene		ND*
2.3.7.8-tetrachlorodibenzo-p-dioxin		ND*
		ND**

ND = < 10,000 mg/kg ND* = < 40,000 mg/kg ND** = < 100,000 mg/kg

^aUnder the conditions stipulated by EPA Method 625 this compound pair cannot be chromatographically resolved. The concentration reported reflects the



Project	5-668-073

GCA Control No. 42240

DATA REPORT SHEET

Acid Extractables

Analysis Date 1/29/85				
Instrument HP 5985 GC/MS				
Ion Used To Quantitate	Concentration (mg/kg)			
_	ND			
	ND**			
	Instrument Ion Used To			

ND = < 10,000 mg/kgND ** = < 100,000 mg/kg



Projec	t	5-668-073
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GCA	Control	No.	42240
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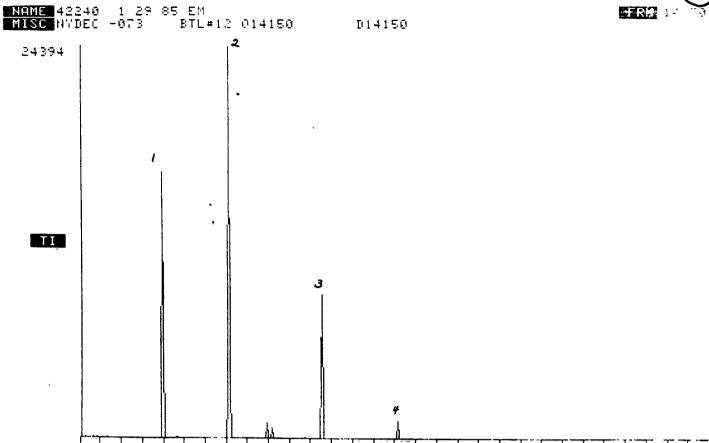
DATA REPORT SHEET

Extractables Qualitative Analysis

Sample I.D. R=684-045-02		Analysis Dat	e 1/29/85	
Sample Matrix Tar	Instrument	HP 5985 GC/MS		
Compound		CAS No.	Concentration* (mg/kg)	
2-Methylnaphthalene		91-57-6	10,000	
		•		

^{*}Concentration based on a comparison of the total ion area of the compound with that of the internal standard.





- $1 = D_4 \div 1, 4$ -dichlorobenzene
- 2 = Dg-naphthalene
- $3 = D_{10}$ -acenaphthene
- $4 = D_{10}$ -phenanthrene

HARBOR POINT PROPERTY: THE MOHAWK RIVER
RESULTS OF INITIAL RIVER SURVEY
STEP 2 RIVER REPORT

This Privileged and Confidential Material is on file at the offices of NYSDEC 50 Wolf Road, Albany, New York

rally occurring nonessential amino acid. The common form is L(+)-aspartic acid. Nontoxic.

Properties: Colorless crystals; soluble in water, insol-

uble in alcohol and ether; optically active.

DL-aspartic acid: M.p. 278-280°C with decomposition; sp. gr. 1.663 (12/12°C).

L(+)-aspartic acid: M.p. 251°C.

D(-)-aspartic acid: M.p. 269-271°C with decomposi-

tion; sp. gr. 1.6613.
Source: Young sugar cane; sugar beet molasses.
Derivation: Hydrolysis of protein; reaction of ammonia with diethyl fumarate.

Uses Biological and clinical studies; preparation of culture media; organic intermediate; dietary supplement; detergents; fungicides; germicides; metal complexation; synthetic sweetener base (L-form).

Available commercially as D(-)-, L(+)-, and DLaspartic acid.

aspartocin. USAN for antibiotic produced by Streptomyces griseus.

aspergillic acid C₁₂H₂₀N₂O₂. 2-Hydroxy-3-isobutyl-6-

(1-methylpropyl)pyrazine 1-oxide. An antibiotic from strains of Aspergillus flavus. Nontoxic.

Properties: Solid. M.p. 97°C; insoluble in cold water, soluble in common organic solvents and dilute acids. Hydrochloride melts at 178°C and is soluble in water.

asphalt (petroleum asphalt, Trinidad pitch, mineral pitch). A dark-brown to black cementitious material, solid or semisolid in consistency, in which the predominating constituents are bitumens, which occur in nature as such or are obtained as residua in petroleum refining (ASTM). It is a mixture of paraffinic and aromatic hydrocarbons and heterocyclic com-

pounds containing sulfur, nitrogen, and oxygen.
Properties: Black solid or viscous liquid; sp. gr. about 1.0; soluble in carbon disulfide. Flash point 450°F; autoignition temp. 900°F; solid softens to viscous liquid at about 200°F; penetration value (paving) 40-300 (roofing) 10-40. Good electrical resistivity. Combustible; low toxicity.

Occurrence: California, Trinidad, Venezuela, Cuba, Canada (Athabassa sa canda)

Canada (Athabasca tar sands).

Containers: Drums, barrels, tank trucks, tank cars. Uses: Paving and road-coating; roofing; sealing and joint filling; special paints; adhesive in electrical laminates and hot-melt compositions; diluent in low-grade rubber products; fluid loss control in hydraulic fracturing of oil wells; medium for radioactive waste disposal; pipeline and underground cable coating; rustpreventive hot-dip coatings; base for synthetic turf; water-retaining barrier for sandy soils; supporter of rapid bacterial growth in converting petroleum components to protein.

See also bacteria; protein; asphalt, blown.

asphalt, blown (mineral rubber, oxidized asphalt, condensed asphalt). Black, friable solid obtained by blowing air at high temperature through petroleum-derived asphalt, with subsequent 'cooling. Penetration value 10-40; softening point 185-250°F. Combustible. Uses are primarily roofing, as diluent in low-grade rubber products, and as thickener in oil-based drilling fluids. Shipped in 55-2al metal drums. based drilling fluids. Shipped in 55-gal. metal drums. For further information on asphalt, refer to the Ashalt Institute, 1270 Avenue of the Americas, New York, N.Y.

asphalt, cut-back. A liquid petroleum product, produced by fluxing an asphaltic base with suitable dis-tillates. (A.S.T.M.)

Properties: Flash point (open cup) 50°F. Solubility

of residue from distillation in carbon tetrachloride 99.5%.

Hazard: Flammable, dangerous fire hazard. Use: Road surfaces.

Shipping regulations: (Rail) Red label. (Air) Flammable Liquid label.

asphaltene. A component of the bitumen in petroleums, petroleum products, malthas, asphalt cements, and solid native bitumens, soluble in carbon disulfide but insoluble in paraffin naphthas. (A.S.T.M.)

asphalt, liquid. See residual oil; asphalt.

asphalt, oxidized. See asphalt, blown.

asphalt paint. Asphaltic base in a volatile solvent with or without drying oils, resisns, fillers, and pigments. Ground asbestos is frequently used as a component of heavy asphaltic paints for roofing and waterproofing purposes.

Hazard: Flammable; dangerous fire risk.

asphyxiant gas. A gas which has little or no positive toxic effect but which can bring about unconsciousness and death by replacing air and thus depriving an organism of oxygen. Among the so-called asphyxiant gases are carbon dioxide, nitrogen, helium, methane, and other hydrocarbon gases.

aspidospermine C22 H30O2N2.

Properties: White to brownish-yellow crystalline alkaloid. M.p. 108°C; b.p. 200°C (I to 2 mm); sublimes at 180°C under reduced pressure. Soluble in fats and fixed oils; sparingly soluble in absolute alcohol and ether. Its sulfate and hydrochloride are soluble in water

Hazard: Moderately toxic by ingestion.

Use: Medicine.

aspirin (acetylsalicylic acid; ortho-acetoxybenzoic acid).

CH₃COOC₆ H₂COOH.

Properties: White crystals or white, crystalline powder. Odorless; slightly bitter taste. Stable in dry air; slowly hydrolyzes in moist air to salicylic and acetic acids. Soluble in water, alcohol, chloroform, and ether; less soluble in absolute ether. Dissolves with decomposition in solutions of alkali hydroxides and carbonates. M.p. 132-136°C; b.p. 140°C (dec.). Derivation: Action of acetic anhydride on salicylic acid.

Method of purification: Crystallization. Grades: Technical, U.S.P.

Containers: 25-lb boxes; 25-, 100-, 250-lb drums. Hazard: An allergen; may cause local bleeding, especially of the gums; 10-gram dosage may be fatal. Dust dispersed in air is serious explosion risk. Use: Medicine (anodyne).

"Aspon." Trademark for a concentrate of tetra-npropyl dithionopyrophosphate, a liquid insecticide.

assay. Determination of the content of a specific component of a mixture, with no evaluation of other components. Such determinations are made on ores of various metals (especially precious metals), on pharmaceutical products to validate the amount of drug present in a given unit, and on organisms (bacteria) to determine their reactions to an antibiotic or insecticide. The latter procedure is called bioassay. Ores are assayed by heat fractionation, organic materials by solvent extraction and chemical separation.

assistant. A term loosely used in the textile industry for any chemical compound that aids in a processing step, e.g., scouring, dyeing, bleaching, finishing, etc. See also auxiliary; dyeing assistant.



"Bentolite."471 Trademark for a series of white bentonites from Texas.

Grades: H and L, high and low gelling.

Uses: Suspending agent; thixotropic agent; adhesives; ceramic bonding and plasticizing agent; desiccant; medicated powders.

"Bentone."304 Trademark for organic derivatives of hydrous magnesium aluminum silicate minerals. Uses: Gelling and pigment-suspending agents.

bentonite. A colloidal clay (aluminum silicate) composed chiefly of montmorillonite. There are two varieties: (1) sodium bentonite (Wyoming or western), which has high swelling capacity in water; and (2) calcium bentonite (southern), with negligible

swelling capacity.
Properties: (Wyoming) Light to cream-colored impalpable powder, forms colloidal suspension in water, with strongly thixotropic properties. Nontoxic; non-

combustible. Occurrence: Wyoming; Mississippi; Texas; Canada;

Italy; U.S.S.R.

Containers: Paper bags; drums; bulk carloads. Uses: Oil-well drilling fluids; cement slurries for oilwell casings; bonding agent in foundry sands and pelletizing of iron ores; sealant for canal walls; thickener in lubricating greases and fireproofing compositions; cosmetics; decolorizing agent; filler in ceramics, refractories, paper coatings; asphalt modifier; polishes and abrasives; food additive; catalyst sup-

port. See also clay.

"Benzahex." 147 Trademark for a group of insecticides containing gamma-benzene hexachloride. Hazard: See benzene hexachloride.

benzalacetone. See benzylidene acetone.

benzalazine (benzylidene azine) C6H3CH:NN:CHC6H5. Properties: Yellow crystals; m.p. 91-93°C; insoluble in cold water; soluble in benzene and hot alcohol. Uses: Stabilizer; polymerization catalyst; ultraviolet absorbent; reagent and intermediate.

benzal chloride. See benzyl dichloride.

benzaldehyde (benzoic aldehyde; synthetic oil of bitter

almond) C6H5CHO.

Properties: Colorless or yellowish, strongly refractive, volatile oil with odor resembling oil of bitter almond, and burning aromatic taste; oxidizes readily; miscible with alcohol, ether, fixed and volatile oils; slightly soluble in water. Sp. gr. 1.0415 (25/4°C; refractive index (20°C) 1.5440-1.5464; f.p. -56°C; b.p. 178°C. Flash point 145°F (C.C.). Oxidizes in air to benzoic acid. Combustible. Autoignition temp. 377°F. Derivation: (a) Air oxidation of toluene with uranium or molybdenum oxides as catalysts; (b) chlorination of toluene with hydrolysis by acid or alkali.

Impurities: Usually chlorine derivatives.

Method of purification: Rectification. Grades: Technical; N.F.. Note: The specifications, especially regarding impurities, vary considerably for the grades used for dye manufacture from those used in perfumery.

Containers: Tins; carboys; drums; tank cars.

Hazard: Moderately toxic by ingestion.

Uses: Chemical intermediate for dyes, flavoring materials, perfumes, and aromatic alcohols; solvent for oils, resins, some cellulose ethers, cellulose acetate and nitrate; flavoring compounds; synthetic perfumes; manufacture of cinnamic acid, benzoic acid; pharmaceuticals; photographic chemicals. See also oil of bitter almond.

benzaldehyde cyanohydrin. See mandelonitrile.

benzaldehyde green. See malachite green.

benzalkonium chloride. A mixture of alkyl dimethylbenzylammonium chlorides of general formula C6H3CH2N(CH3)2RCI in which R is a mixture of the alkyls from C₈H₁₇ to C₁₈H₃₇. It is a typical quaternary ammonium salt.

Properties: White or yellowish-white, amorphous powder or gelatinous pieces. Aromatic odor and very bitter taste; soluble in water, alcohol or acetone; almost insoluble in ether; slightly soluble in benzene. Water solutions foam strongly when shaken and are alkaline to litmus.

Grade: U.S.P.

Hazard: Toxic by ingestion and skin absorption Uses: Cationic detergent; surface antiseptic, fungicide.

benzamide (benzoylamide) C₆H₃CONH₂

Properties: Colorless crystals; m.p. 130°C, b.p. 288°C; sp. gr. 1.341. Soluble in hot water, hot benzene, alcohol, and ether. Combustible.

Derivation: From benzoyl chloride and ammonia or ammonium carbonate.

Grades: Technical

Uses: Organic synthesis.

benzaminoacetic acid. See hippuric acid.

benzanilide (benzoylaniline; phenylbenzamide)

C₆H₅NH(COC₆H₅).
Properties: White to reddish crystals and powder, closely related to acetanilide, containing benzoyl in place of acetyl radical. Sp. gr. 1.306; m.p. 160-162° C Soluble in alcohol; insoluble in water; slightly soluble in ether.

Derivation: From benzoic anhydride and aniline with caustic soda.

Uses: Intermediate in the synthesis of dyes, drugs and perfumes.

benzanthrone C17H10H, a four-ring system.

Properties: Pale yellow needles; soluble in alcohol and other organic solvents. M.p. 170°C

Derivation: (a) From anthranol and glycerol by condensation by means of sulfuric acid (anthranol is made from anthraquinone); (b) from anthracene in sulfuric acid solution by addition of glycerol and heating to 100-110°C until the anthracene disappears. The reaction mass is then diluted with water, salted out and purified.

Method of purification: Crystallization from toluene. Use: Dves.

benzathine penicillin G (N,N1-dibenzylethylenediamine dipenicillin G) 2C₁₆H₁₆N₂O₄S· C₁₆H₂₀N₂· 4H₂O.
roperties: White, odorless, crystalline powder;

Properties: slightly soluble in alcohol; practically insoluble in water; pH of a saturated solution is 4.5-7.5. Grade: U.S.P.

Use: Medicine (antibiotic).

benzazimide. See 4-ketobenzotriazine.

"Benzedrine."71 Trademark for amphetamine sulfate.

benzene C6H6. Thirteenth in order of high-volume chemicals produced in U.S. (1975).

mandelonitrile. nali nite green.

mixture of alkyl dimethylorides of general formula h which R is a mixture of the n w t is a typical quaternary

yellowish-white, amorphous eces. Aromatic odor and very water alcohol or acetone; al-; sle itly soluble in benzene, trongry when shaken and are

d skin absorption. ce antiseptic; fungicide. ion su:

) C₆H₅CONH₂. stals; m.p. 130°C; b.p. 288°C; in het water, hot benzene, al-oust the

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eddish crystals and powder, anilia, containing benzoyl in Sp. 1.306; m.p. 160-162°C. Solume in water; slightly sol-

ic anhydride and aniline with hesis of dyes, drugs and e sy

four-ring system. meedles; soluble in alcohol and M. 170°C.

nthre ol and glycerol by control of surruric acid (anthranol is

one); (b) from anthracene in by addition of glycerol and un the anthracene disap-ass in hen diluted with water, 155

Crystallization from toluene.

inbenzylethylenediamine · C₁₆H₂₀N₂ · 4H₂O. N_1N_1 iorless, crystalline powder; ohol; practically insoluble in d salution is 4.5-7.5.

c). enzotriazine.

rk fas amphetamine sulfate. order of high-volume

Structure: I. Complete ring showing all elements. II. Standard ring showing double bonds only.

III. Simple ring without double bonds, with numerals indicating position of carbon atoms to which substituent atoms or groups may be attached (2 = ortho, 3 = meta, 4 = para).

IV. Generalized structure, with enclosed circle suggesting the resonance of this compound. These structures are also referred to as the benzene

Properties: Colorless to light-yellow, mobile, nonpolar hquid of highly refractive nature; aromatic odor; vapors burn with smoky flame; b.p. 80.1°C; m.p. 5.5°C; sp. gr. 0.8790 (20/4°C); wt/gal 7.32 lb; refractive index (n 20/D) 1.50110; flash point (closed cup) 12°F; surface tension 29 dynes/cm. Autoignition temp. 1044° F. Miscible with alcohol, ether, acetone, carbon tetrachloride, carbon disulfide, acetic acid; slightly soluble in water.

Derivation: (a) Hydrodealkylation of toluene or of pyrolysis gasoline (q.v.); (b) transalkylation of toluene by disproportionation reaction; (c) catalytic reforming of petroleum; (d) fractional distillation of

Grades: Crude; straw color; motor; industrial pure (2°C); nitration (1°C); throphene-free; 99 mole %; 99.94 mole %; nanograde.

Containers: Drums; tank cars; barges. Hazard: Flammable, dangerous fire risk. Explosive limits in air, 1.5 to 8% by volume. Toxic by ingestion, inhalation, and skin absorption. Tolerance, 25 ppm in air. Safety data sheet available from Manufacturing Chemists Assn., Washington, D.C.

Uses: Ethylbenzene (for styrene monomer); dodecylbenzene (for detergents); cyclohexane (for nylon); phenol; nitrobenzene (for aniline); maleic anhydride; dodecylbenzene; chlorobenzene; diphenyl; benzene hexachloride; benzene-sulfonic acid; solvent; antiknock gasoline.

Shipping regulations: (Rail) Red label. (Air) Flam-mable Liquid label. See also aromatic.

benzene azimide. See 1,2,3-benzotriazole.

benzeneazoanilide. See diazoaminobenzene.

benzeneazobenzene. See azobenzene.

benzeneazo-para-benzeneazo-beta-naphthol III, tetraazobenzene-beta-naphthol) C₆H₅NNC₆H₄-4-NN-1-C₁₀H₆-2-OH. A red dye; C.I. 26100. Properties: Brown powder; m.p. 195°C; insoluble in

water; soluble in alcohol and oils. Uses: Coloring oils red; biological stain.

benzeneazonaphthylethylenediamine. See azodine.

benzenecarboxylic acid. See benzoic acid.

benzenediazonium chloride C6H5N(N)Cl. Properties: Ionic salt. Very soluble in water; insoluble in most organic solvents.

Hazard: Explodes on heating.

Use: Dye intermediate.

Shipping regulations: Not listed. Consult authorities.

benzene dibromide. See dibromobenzene

benzene-ortho-dicarboxylic acid. See phthalic acid.

benzene-para-dicarboxylic acid. See terephthalic acid.

benzene hexachloride (BHC). A commercial mixture of isomers of 1,2,3,4,5,6-hexachlorocyclohexane (q.v.), insecticide. The gamma isomer is toxic. Use may be restricted. See also lindane.

benzenemonosulfonic acid. See benzenesulfonic acid.

benzenephosphinic acid (phenylphosphinic acid) CaHaHaPOs.

Properties: Colorless crystals; m.p. 82-84°C; sp. gr. 1.376 (29°C). Decomposes at 200°C. Stable in air. Soluble in water, alcohol, acetone. Slightly soluble in ether; insoluble in benzene, hexane, carbon tetrachloride. Combustible.

Containers: 100-lb fiber drums

Uses: Antioxidant; intermediate for metallic salt formation; accelerator for organic peroxide catalysts.

benzenephosphonic acid (phenylphosphonic acid) C₆H₅H₂PO₃.

Properties: Colorless crystals. M.p. 158°C; sp. gr. 1.475 (4°C); decomposes at 275°C; soluble in water, alcohol, carbon tetrachloride. Combustible.

Containers: 100-lb fiber drums.

Uses: Intermediate in antifouling paint agents; catalyst in organic reactions.

benzenephosphorus dichloride C₆H₅PCl₂.

Properties: Highly reactive colorless liquid. M.p. -51°C; b.p. 224.6°C; sp. gr. 1.315 (25°C); refractive index 1.5958 (n 25/D). Soluble in common inert organic solvents; fumes in air; hydrolyzes in water. Containers: 55-gal stainless steel drums

Hazard: Flammable; corrosive to skin and tissue. Uses: Organic synthesis, for derivation of plasticizers,

polymers, antioxidants: oil additives. Shipping regulations: (Rail) Red label. (Air) Corrosive label. Not acceptable on passenger planes.

benzenephosphorus oxydichloride C6H5POCl2 Properties: Reactive colorless liquid. M.p. 3.0°C; b.p. 258°C; sp. gr. 1.197 (25°C); refractive index 1.5585 (n 25/D). Soluble in common inert organic solvents; hydrolyzes in water. Combustible.

Containers: 55-gal nickel drums. Hazard: Strong irritant to skin.

Uses: Organic synthesis, for derivation of plasticizers, polymers, antioxidants, oil additives.

benzenesulfonic acid (benzenemonosulfonic acid; phenylsulfonic acid) C₆H₃SO₃H.

Properties: Fine, deliquescent needles or large plates;

m.p. 65-66°C when anhydrous; with 1.5 molecules water, m.p. is 43-44°C; soluble in water and alcohol; slightly soluble in benzene; insoluble in ether and carbon disulfide.

Derivation: By reacting benzene with fuming sulfuric acid.

Uses: Manufacture of phenol, resorcinol and other organic syntheses, and as a catalyst.

benzene-1,3-5-tricarboxylic acid chloride. See trimesoyl trichloride.

benzenoid. Any organic compound containing or derived from the benzene ring structure, e.g., phenol,



"Coagulant Aid." 108 Trademark for a series of polyelectrolytes and combinations of polyelectrolytes with other materials.

Uses: Clarification of water for municipal and industrial uses

coagulation. Irreversible combination or aggregation of semisolid particles such as fats or proteins to form a clot or mass. This can often be brought about by addition of appropriate electrolytes, for example, by the addition of an acid to milk or of aluminum sulfate to turbid water. Mechanical agitation and removal of stabilizing ions, as in dialysis, also cause coagulation. The clotting of blood by thrombin (q.v.), the coagulation of rubber particles in latex by acetic acid and of egg-white by heat are additional instances. See also flocculation.

Coahran process. Recovery of acetic acid from pyroligneous acid by extracting with ether It is an im-proved version of the Brewster process (q.v.), but is basically the same

A natural solid combustible material consisting chiefly of amorphous elemental carbon with low percentages of hydrocarbons, complex organic compounds and inorganic materials. Coal was formed from prehistoric plant life and occurs in layers or veins in sedimentary rocks. It is far more plentiful in the U.S. than petroleum and is an important source of heat and energy; its use as a fuel is increasing. In the U.S. it occurs chiefly in West Virginia and Kentucky, as well as in Wyoming and other western states. Much of it is too high in sulfur content to meet present pollution standards.

Coal is classified according to its heating value, expressed in Btu/lb, and its fixed carbon content.

Anthracite	Fixed Carbon	Btu/lb
Low to Medium	86-98%	13,500-15,600
Volatile Bituminous High Volatile	69-86%	14,500-15,600
Bituminous	46-69%	11,000=15,000
Subbituminous	46-60%	8,300=13,000
Lignite	46-60%	5,500= 8,300

Coal is most frequently specified in terms of its proximate analysis, giving the percentages of moisture, volatile combustible matter, fixed carbon, and ash. An ultimate analysis gives the percentages of the various elements present (C, H, O, N, and S).

Coal is also an important source of chemical raw

materials: pyrolysis (destructive distillation) yields coal tar and hydrocarbon gases, which can be upgraded by hydrogenation or methanation to synthetic crude oil and fuel gas, respectively; catalytic hydrogenation yields hydrocarbon oils and gasoline; gasi-fication produces carbon monoxide and hydrogen (synthesis gas) from which ammonia and other products can be made. Numerous processes for adapting these reactions to large-scale production of fuel oil and gasoline are in the pilot-plant stage, though none has yet been commercially developed. Methane is being produced from coal mines on a commercial scale.

See also gasification; hydrogenolysis; hydrosolvation. Note: Pending the further development of alternative energy sources, coal is probably the safest and most abundant fuel available in the U.S. A large percentage can be obtained by surface or strip mining, though the necessary restoration of the environment (soil and plant life) is expensive. Improvements in safety are needed in order to realize full and safe production from deep mines.

coal gas (bench gas; coke-oven gas) A mixture of gases produced by the destructive distillation of bituminous coal in highly heated fire-clay or silica retorts or in by-product coke ovens.

Hazard: Flammable; highly toxic. Uses: Directly in open hearth furnaces.

Shipping regulations: (Rail) Red Gas label. (Air) Flammable Gas label. Not accepted on passenger

coalescence. The union of two or more droplets of a liquid to form a larger droplet, brought about when the droplets approach one another closely enough to overcome their individual surface tensions. The combination of fine mercury droplets is an example Coalescing agents are used to remove solid con-taminants from hydrocarbons; coalescence may also be brought about by mechanical means in centrifuges

coal, gasification. See gasification.

coal, hydrogenation. See hydrogenolysis: gasification

coal oil. The crude oil obtained by the destructive distillation of bituminous coal; or the distillate obtained from this oil.

Hazard: Flammable, moderate fire risk.

Properties: Black, viscous liquid (or semi-solid). naphthalene-like odor; sharp burning taste; obtained by destructive distillation of bituminous coal, as in cokeovens; one ton of coal yields 8.8 gallons of coal tar. Combustible. Sp. gr. 1.18 to 1.23 (60 60°F). Soluble in ether, benzene, carbon disulfide, chloroform; partially soluble in alcohol, acetone, methanol, and benzene; only slightly soluble in water.

Coal tar may be hydrogenated under pressure to form a petroleum-like fuel suitable for residual use. Coal-tar fractions obtained by distillation and the chemicals found in each are as follows: (1) light oil (up to 200°C); benzene, toluene, xylenes, cumenes, coumarone, indene; (2) middle oil (200-250°C) and (3) heavy oil (250-300°C): naphthalene, acenaphthene, methylnaphthalenes, fluorene, phenol, cresols, pyridine, picolines; (4) anthracene oil (300-350°C) phenanthrene, anthracene, carbazole, quinolines; and (5) pitch (q.v.). Typical yields: 5% light oil; 17% middle oil; 7% heavy oil; 9% anthracene oil; 62% pitch. Treatment with alkalies, acids, and solvents is necessary to separate the individual chemicals. Grades: Crude, refined; U.S.P.

Containers: Tank cars; barrels. Hazard: Toxic by inhalation.

Uses: Raw material for plastics, solvents, dyes, drugs, and other organic chemicals. The crude or refined product or fractions therof are also used for waterproofing, paints, pipecoating, roads, roofing, insulation, as pesticides and sealants, and in medicine.

coal-tar distillate. The lighter fractions of coal tar.
The terms coal tar light oil, coal tar naphtha, and coal tar oil are loosely defined and are sometimes

regarded as synonymous.

Hazard: Flammable, dangerous fire risk. Toxic by inhalation and skin absorption.

Shipping regulations: (Rail) Red label. (Air) Flammable Liquid label.

See also entries under naphtha.

coal-tar dye. A dye produced from the coal-tar hydrocarbons or their derivatives such as benzene, toluene, xylene, naphthalene, anthracene, aniline, etc. See also dye, synthetic.

Shipping regulations: (Air) Coal-tar dye, liquid, n.o.s., Corrosive label.

"CRESTAPOL"

r 1,2,3,4-cyclopentanecarrk

rk for allyl diglycol carbonate.

um.

tica plastic highly resistant to in. Trinished as a clear liquid.

nses, shields, instrument panels. gla

oosi on by heat, with or without oleum or heavy petroleum frac-on of gasoline, fuel oil, or other Steel cracking of naphtha to personnel is widely prac-also applied to other thermal esses; thus ammonia (NH₃) may ogen and hydrogen, and natural ach methane (CH4) are cracked hydrogen, or into other hydro-

ng, reforming, catalysis.

f agricultural chemicals in-

(active ingredient, butoxy poly-coloriess liquid, 100% active ma-

(active ingredient, 3,5-dimethyl-this azine-2-thione). Wettable margial. Toxic and irritant. on (active ingredient, 2-heptadecyl 34% active solution.

SES) (active ingredient, sodium yeth sulfate). Water-soluble material. Tolerance, 10 mg per

vegetable oil obtained from the a plant related to mustard and proper sing techniques have been grice ural Research Service of can be grown on marginal and s high content of erucic acid and ontent of its meal make it eco-; the oil itself is useful as an in-especially for molds for contin-

otassium bitartrate.

N-g nylgiycine, (alpha-methyl-) H C(NH₂)N(CH₃)CH₂COOH. I wuely distributed in the musody.

drate) Prisms from water; anhycon oses 303°C; slightly soluble

in earr. ly isolated from meat extracts. ..P.

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nhydride of creatine (q.v.);

from water; decomposes about water; slightly soluble in alcohol; acetone, and chloroform. ether ear

-methylphenol; 4-methylguaiacol) H. One of the active constituProperties: Colorless to yellow liquid. Sp gr. 1.092 (25°C); m.p. 5.5°C; b.p. 220°C; slightly soluble in water; soluble in alcohol, benzene, chloroform, ether, acetic acid.

creosote carbonate.

Properties: Clear, colorless or yellowish, viscous liquid; slight creosote odor and taste. Soluble in alcohol; insoluble in water.

Derivation: Mixture of carbonates of various constituents of creosote. Use: Medicine.

creosote, coal-tar (creosote oil; liquid pitch oil; tar oil). Properties: Yellowish to dark green-brown, oily liq-uid; clear at 38°C or higher; naphthenic odor; frequently contains substantial amounts of naphthalene and anthracene; distilling range 200-400°C; flash point 165°F (closed cup); soluble in alcohol, benzene, and toluene; immiscible with water Combustible.

Derivation: Fractional distillation of coal-tar.

Method of purification: Rectification.

Grades: Technical; crude; refined.

Containers: Drums; tank cars. Hazard: Toxic by inhalation of fumes; skin and eye

irritant. Use may be restricted.
Uses: Wood preservative (ties, telephone poles, marine piling, etc.); disinfectants.

para-cresidine. See 5-methyl-ortho-anisidine.

"Creslan."57 Trademark for an acrylic fiber.

cresol (methyl phenol; hydroxymethylbenzene) CH₂C₆H₄OH. A mixture of isomers obtained from

coal tar or petroleum.

Properties: Colorless, yellowish, or pinkish liquid; phenolic odor; sp. gr. 1.030-1.047; wt/gal 8 66-8.68 lb; flash point approx. 180°F; m.p. 11-35°C; b.p. 191-203°C. Soluble in alcohol, glycol, and dilute alkalies. Combustible.

Derivation: Coal tar (from coke and gas works), also

synthetic.

Grades: Various, depending on phenol content, or other properties. N.F. grade contains not more than 5% phenol.

Containers: Drums; tank cars; tank trucks.

Hazard: Toxic and irritant; corrosive to skin and mucous membranes; absorbed through skin. Tolerance, 5 ppm in air. Safety data sheet available from Manufacturing Chemists Assn., Washington, D.C.

Uses: Disinfectant; phenolic resins; tricresyl phosphate; ore flotation; textile scouring agent; organic intermediate; mfg. of salicylaldehyde, coumarin, and herbicides; surfactant; synthetic food flavors (para isomer only).

Shipping regulations: (Air) Poison label.

See also cresylic acids.

meta-cresol (meta-cresylic acid; 3-methylphenol) CH1C6H4OH.

Properties: Colorless to yellowish liquid; phenol-like odor. Soluble in alcohol, ether, and chloroform; soluble in water. Sp. gr. 1.034; m.p. 12°C; b.p. 203°C; wt/gal 8.66 lb flash point 187°F. Combustible. Autoignition temp. 1038°F.

Derivation: By fractional distillation of crude cresol (from coal tar); also synthetically.

Method of purification: Rectification. Grade: Technical (95-98%).

Toxicity, uses, see cresol.

ortho-cresol (ortho-cresylic acid; 2-methylphenol)

CH3C6HLOH.

Properties: White crystals; phenol-like odor. Soluble in alcohol, ether, chloroform, and hot water. Sp. gr. 1.047; m.p. 30.9°C; flash point 178°F. Autoignition temp. 1110°F; b.p. 191°C; lb/gal 8.68. Combustible. Derivation: (a) By fractional distillation of crude cresol from coal tar. (b) Interaction of methanol and

Method of purification: Crystallization. Grades: According to freezing point: 25°, 29°, 30°, 30.5°C, etc.

Toxicity, uses, see cresol.

para-cresol (para-cresylic acid; 4-methylphenol) CH₃C₆H₄OH.

Properties: Crystalline mass; phenol-like odor. Soluble in alcohol, ether, chloroform, and hot water, wt/gal 8.67 lb. Sp. gr. 1.039; b.p. 202°C; m.p. 35.26°C; flash point 187°F. Combustible. Autoignition temp. 1038°F.

Derivation: (a) By fractional distillation of crude cressel (b) From heavens by the cumere process (see

sol. (b) From benzene by the cumene process (see

Method of purification: Crystallization Grades: Technical; 98%; 99.0% min purity or 34°C min F.P.

Toxicity, uses, see cresol.

cresolphthalein C6H4COOC(C6H3(OH)CH3)2. An acid-

base indicator, changes from colorless to red between pH 8.2 and 9.8. See also indicator.

cresol purple C6H4SO2OC(C6H3(OH)CH3)2. Meta-

cresolsulfonphthalein, an acid-base indicator, showing color change from red to yellow over the range pH 1.2 to 2.8 and from yellow to purple over the range pH 7.4 to 9.0. See also indicator.

cresol red CoH4SO2OC(CoH1(OH)CH3)2. Ortho-cresol-

sulfonphthalein, an acid-base indicator, changes from red to yellow between pH 0.2 and 1.8 and from yellow to red between pH 7.0 and 8.8. See also indicator.

cresotic acid (cresotinic acid; hydroxytoluic acid) CH₃C₆H₃(OH)COOH. Ten possible isomers; most common is 2-hydroxy-3-methylbenzoic acid, also known as ortho-cresotic acid or ortho-homosalicylic acid. The description which follows is of this isomer.

Properties: White crystals or powder; m.p. 166°C; b.p. about 250°C; insoluble in water; soluble in alcohol and ether. Combustible.

Derivation: Treatment of ortho-cresol with caustic and carbon dioxide under pressure.

Containers: Fiber cans; drums.

Uses: Dye intermediate; research on plant growth inhibition.

"Crestalkyd."²⁶³ Trademark for a group of oil-modified alkyd resins. Oil length from 30% to 80%, A.V.

Uses: High-quality paints, lacquers, enamels, and varnishes. Plasticizing resin for nitrocellulose and similar finishes.

"Crestapol."263 Trademark for a series of plasticizers for polyvinyl chloride; also a designation for specialty polyesters used in dispersing media.

Superior numbers refer to Manufacturers of Trade Mark Products. For page number see Contents.

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i,

(c) cleaners'. A dry-cleaning fluid derived from petroleum and similar to Stoddard solvent (q.v.), but not necessarily meeting all its specifications. Flash point 100°F.

(2) coal-tar (a) heavy (high-flash naphtha)

Properties: Deep amber to dark red liquid; a mixture of xylene and higher homologs; sp. gr. 0.885-0.970; b.p. 160-220°C (about 90% at 200°C); flash point 100° F; evaporation 303 minutes.

Derivation: From coal-tar by fractional distillation.

Containers: Drums; tank cars.

Hazard: Toxic by ingestion, inhalation, and skin absorption. Moderate fire risk. Tolerance, 100 ppm in air.

Uses: Coumarone resins; solvent for asphalts, road Uses: Coumarone resins; solvent for aspitants, road tars, pitches, etc.; cleansing compositions; process engraving and lithography; rubber cements (solvent); naphtha soaps; mfg. of ethylene and acetic acid.

(b) solvent (160° benzol).

Properties: A mixture of small percentages of benzene and toluene with xylene and higher homologs from coalder. (a) Crude: dark straw-colored liquid.

from coal-tar. (a) Crude: dark straw-colored liquid.
(b) Refined: water-white liquid. Sp. gr. (a) 0.862-0.892, (b) 0.862=0.872; b.p. (a) about 160°C (80%), (b) about 160°C (90%); flash point (a) and (b) about

Derivation: From coal-tar by fractional distillation.

Grades: Dark straw, water-white. Containers: Drums; tank cars. Hazard: Flammable, dangerous fire risk.

Uses: Solvent; xylene; cumene; nitrated, for incorporation in dynamite.

Shipping regulations: (Rail) Red label. (Air) Flammable Liquid label.

naphthacene (tetracene; rubene) C18H12. The molecule

consists of four fused benzene rings.

Properties: Orange solid; sp. gr. 1.35; m.p. about 350°C; not easily soluble; slight green fluorescence in daylight.

Occurrence: In commercial anthracene and coal tar. Hazard: Explodes when shocked; reacts with oxidizing materials.

Uses: Organic synthesis.

naphthalene (tar camphor) C₁₀H₈.



Properties: White crystalline, volatile flakes; strong coal-tar odor. Soluble in benzene, absolute alcohol and ether; insoluble in water. Sp. gr. 1.145 (20/4°C); m.p. 80.2°C; b.p. 217.96°C; flash point 176°F. Sublimes at room temperature. Autoignition temp. 979°F. Combustible.

Derivation: (a) From coal-tar oils boiling between 200 to 250°C (middle oil) by crystallization and distillation. (b) From petroleum fractions after various

catalytic processing operations.

Grades: By melting point, 74°C min (crude) to above 79°C (refined); scintillation (80-81°C).

Forms: Flakes, cubes, spheres, powder.

Containers: Cans; drums; tank cars. Hazard: Toxic by inhalation. Safety data sheet available from Manufacturing Chemists Assn., Washington, D.C. Tolerance, 10 ppm in air.
Uses: Intermediate (phthalic anhydride, naphthol,

"Tertralin," "Decalin," chlorinated naphthalenes, naphthyl and naphthol derivatives, dyes); moth repellent; fungicide; explosives; cutting fluid; lubricant; synthetic resins; synthetic tanning; preservative; solvent; textile chemicals; emulsion breakers; scintillation counters.

Shipping regulations: (Air) Crude or refined: Flam-mable Solid label.

alpha-naphthaleneacetic acid (1-naphthylacetic acid) C₁₀H₇CH₂COOH.

Properties: White crystals, odorless; m.p. 132=135°C. Soluble in acetone, ether, and chloroform; slightly soluble in water and alcohol.

Grades: Usually supplied in dilute form, either as a powder or liquid solution ready for use.

Containers: Powder; fiber cans or multiwall paper sacks; solution: glass bottles and carboys. Hazard: Moderately toxic; skin irritant.

Uses: Inducing rooting of plant cuttings; spraying apple trees to prevent early drop.

alpha-naphthaleneacetic acid, methyl ester (MENA) C₁₀H₇CH₂COOCH₃. A plant growth regulator or hormone, used for delaying sprouting of potatoes, weed control, thinning of peaches, etc.

naphthalene, chlorinated. See chlorinated naphthalene. naphthalenediamine. See naphthylenediamine.

1,5-naphthalene diisocyanate C10H6(NCO). White to light yellow crystalline solid; m.p. 127-131°C. Hazard: Toxic and irritant. Use: Manufacture of polyurethane solid elastomers.

naphthalene-1,5-disulfonic acid (Armstrong's acid)

C₁₀H₆(SO₃H)₂.
Properties: White crystalline solid; soluble in water. Derivation: Sulfonation of naphthalene with furning sulfuric acid at low temperature and separation from the 1,6 isomer. Combustible. Hazard: May be toxic.

Uses: Intermediate for dyes.

naphthalene-2,7-disulfonic acid C₁₀H₆(SO₁H)₂. Properties: White crystalline solid; soluble in water. Combustible.

Derivation: Sulfonation of naphthalene at high temperature and separation from 2,6-isomer. Hazard: May be toxic. Use: Intermediate for dyes.

alpha-naphthalenesulfonic acid C10H7SO1H · H2O. Properties: Deliquescent crystals; soluble in water, alcohol, and ether. M.p. 90°C. Combustible.
Derivation: Interaction of naphthalene and sulfuric acid.

Hazard: May be toxic. Hazard: May be toxic.

Uses: Starting point in the manufacture of alphanaphthol, alphanaphtholsulfonic acid, alphanaphthylaminesulfonic acid; solvent (sodium salt) for phenol in the manufacture of disinfectant soaps.

beta-naphthalenesulfonic acid C10H7SO1H or C10 H7SO3 H · H2O.

Properties: Non-deliquescent, white plates; m.p. 124-125°C. Soluble in water, alcohol, and ether. Combustible.

Derivation: Sulfonation of naphthalene.

Hazard: May be toxic.

Uses: Starting point in the manufacture of betanaphthol, beta-naphtholsulfonic acid, beta-naphthyl-aminesulfonic acid; etc.

Superior numbers refer to Manufacturers of Trade Mark Products. For page number see Contents.

UNDERGROUND GASIFICATION OF COAL

The United States, Britain, and the Soviet Union have made tests to gasify coal underground to obtain a low-Btu gas. In this approach, input- and output-bore holes are drilled down to a coal bed at selected points, an oxidizing agent is pumped down the input holes, the coal is ignited, and the gaseous combustion products, including H., CO, and CO., are obtained from the output-bore holes. Where air is used as the oxidizing agent, methane forms only about 5% of the output-gas volume. This percentage can be increased, for example, by using oxygen rather than air as the oxidiant. So far, only the Soviet Union is believed to have practiced underground gasification of coal on a commercial basis. However, this technique may prove to be the most practical way to obtain energy from coal in very deep seams and other seams where mining is impracticable.

Further Reading: Office of Coal Research, Evaluation of Coal Gasincation Technology, Part 1—Pipeline Quality Gas (U.S. Department of the Interior 1973); Coal and the Present Energy Situation," Science, vol. 183, no. 4124, Feb. 8, 1974

to a liquid hydrocarbon such as synthetic crude oil (syncrude) or low-sulfur fuel oil. The pioneering work in this field was done by the German chemist Friedrich Bergius, who patented a process for converting coal to oil in 1913, and the German chemists Franz Fischer and Hans Tropsch, who developed another conversion process in 1923. See also Bergius, Friedrich; Fischer-Tropsch Process.

Plants that produced coal-derived liquid fuels were established in Germany, Japan, England, and other countries having poor petroleum resources, but these fuels generally played a temporary minor role because they cost considerably more than petroleum products. In the 1970's, however, petroleum costs and energy shortages led to renewed interest in coal liquefaction.

Processes. In the Fischer-Tropsch process a coal-derived gas consisting of carbon monoxide and hydrogen is passed through a catalytic reactor, yielding liquid hydrocarbon products. This process is in limited use for commercial production of liquid hydrocarbons.

Pilot-plant studies of other processes can be divided into two classes—coal hydrogenation processes, in which coal is reacted with hydrogen; and coal pyrolysis processes, in which coal is decomposed into liquids by heat.

In the Synthoil process developed by the U.S. Bureau of Mines, crushed coal is slurried with a coal-derived oil. The slurry and turbulently flowing hydrogen are fed to a fixed-bed catalytic reactor, in which the coal is liquefied and desulfurized. The yield is about three barrels of low-sulfur fuel oil per ton of coal. Other coal hydrogenation processes include the H-coal process developed by Hydrocarbon Research, Inc.

In the Coed process developed by the FMC Corporation, crushed coal is heated to successively higher temperatures in a series of fluidized-bed reactors. After further processing, the main products are syncrude, fuel gas, and residual char. The syncrude yield is about one barrel per ton of coal. Another coal pyrolysis process is the Oil Shale Corporation's Toscoal process.

Further Reading: Institute of Gas Technology, Clean Fuels from Coal (Institute of Gas Technology 1973).

COAL TAR is a heavy, dark viscous liquid obtained from the destructive distillation, or carbonization of coal. The primary product of the carbonization of coal is coke, while the secondary products are coal tar, light oil, ammonia liquor, and coal gas.

Production. Modern coke ovens are equipped for the efficient recovery of coal tar and the other by-products of the coking process. The overs, which are erected together in batteries, are long narrow brick chambers. The coal is coked by the combustion of a fuel gas in flues in the walk. In the commonly used high-temperature carbonization process, the oven walls are at a temperature of from 870° to 1205° C (1600° to 2200° F). As the coal carbonizes, the tar and other compounds are separated from the generated gas by cooling in a collecting main; the liquid stream passes to a decanter where the tar and ammoniacal liquor form two layers and are separated. The coal tar is then distilled to give light and middle oils, methylnaphthalenes, and three creosote portions. The residue remaining from the distillation is refined coal tar or coal tar pitch. The term pitch is used to designate the semisolid or solid residues of distillation, while tar is used to refer to

the liquid residues. See also Corr.

Composition and Properties. Coal tar is a complex mixture of organic compounds, most of which are aromatic hydrocarbons. Light oil, which is the fraction that distills off at temperatures of up to 200° C (392° F), contains benzene, toluene, xylene, and various tar acids and bases. The middle oil, which distills off at from 200° to 250° C (392° to 482° F), is primarily composed of tar acids and bases and naphthalene. Heavy oils are recovered at temperatures of from 250° to 300° C (482° to 572° F). This fraction includes the methylnaphthalenes and the creosotes, which contain various aromatic compounds such as phenanthrene, chrysene, and pyrene. At temperatures of from 300° to 350° C (572° to 662° F), phenanthrene, anthracene, carbazole, and quinoline distill off, and the residue of this distillation is the coal tar pitch.

Coal tar pitch has been shown to contain approximately 93% carbon, 4.5% hydrogen, and small amounts of nitrogen, oxygen, and sulfur. More than 100 different compounds have been identified in the coal tar pitch. The molecules found in pitch are relatively small, with molecular weights of from 200 to 2,000. Pitch residues usually exhibit a great change in viscosity with changes in temperature. Both coal tars and coal tar pitches have been found to be extremely resistant to water absorption and to moisture

Modification of Fitch. Depending on the degree of reduction by distillation and other treatments, the flow properties and the hardness of the pitch can be varied greatly. Thus, properties can be changed by the addition of oils or solvents or by additional polymerization. Modification of the pitch can also be effected by the addition of coal fines. Upon heating, the fines are "digested" and this process lowers the susceptibility of the pitch to alterations in viscosity due to changes in temperature.

Uses. Coal tar is used in the production of chemically cured thermosetting mixtures. It has been used with epoxies, urethanes, and polysulfides in the preparation of various chemically resistant and antiskid coatings and sealants. Coal tars are used in road construction and mainte-



wance, although the lectrosed since Westernsed use of pare also used on are very resistant form oils. In additional tangent of the partition of the partition of the partition of the lectrose was although the lectrose with the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was although the lectrose was also used on a lectrose was a lectrose was also used on a lectrose was a lectrose

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Pitch is also use and as a binder for twee. Further uses entain types of routurant for fiber prewage; as a coating base in the prepar matings; for protectures against a marin vent corrosion in u as lines.

Creosote, which is of great and lonpreservative for wood ects such as telepholock pilings.

While a great nu recovered from the clively few of these clarge enough quantit importance. Naphth acids, including phen the most important of also recovered from quantities than the o

Naphthalene is ar production of dyes, chemicals. It is also upellents, lubricants, ar tervative and solvent. production of dyes and tive and solvent. The insecticides and wood infectants. Anthracene ture of dyes.

from coal appeared in However, it was not ur cial uses of coal tar we most recovered coal tar In 1881 the Germans s by-product coking over metallurgical grade co these ovens, it became tars and pitch in large c

It was gradually revents and wood preser from coal tars and that for highway pavement to coatings. Extensive use and especially of coal concurrently with the and dyes from the disti

dan viscous liquid obtained listillation, or carbonization, product of the carboniza-white the secondary product of ammonia liquor, and

ry of coal tar and the other coking process. The ovens, get in batteries, are long, rs. The coal is coked by the el gas in flues in the walls, are at a temperature carbon-over walls are at a temperature. C (1600° to 2200° F). zes, the tar and other coming the companing the liquid stream the generated gas by inguinain; the liquid stream the tar and ammoniacal to give light and middle oils, and three creosote portions, and three creosote portions, and three creosote portions, and three creosote portions oallar pitch. The term pitch et the semisolid or solid resiwhile tar is used to refer to See also COKE.

See also COKE.

Precrities. Coal tar is a compress compounds, most of ic hydrocarbons. Light oil, on that distills off at temperature, and various tar acids and oil which distills off at from 392° to 482° F), is primarily cids and bases and naphthater ecovered at temperatures (482° to 572° F). This me methylnaphthalenes and the contain various aromatic compensathrene, chrysene, and pyther of from 300° to 350° C), thenanthrene, anthracene, inoline distill off, and the resistation is the coal tar pitch.

he been shown to contain colon, 4.5% hydrogen, and niebgen, oxygen, and sulfur. Ifferent compounds have been coal tar pitch. The molecules of the colon, with molecules of the colon, with molecules great change in viscosity with the colon found to be extremely extractive. Both coal tars and coal been found to be extremely extractive.

Pitch. Depending on the degree distillation and other treatments, and the hardness of the pitch read. Thus, properties can be addition of oils or solvents or by merization. Modification of the effected by the addition of he ing, the fines are "digested lowers the susceptibility of the ons in viscosity due to changes in

ar is used in the production of difference thing mixtures. It has a exist, urethanes, and polypreparation of various chemically tiskid coatings and sealants. Coal or read construction and maintenance, although their use for these purposes has decreased since World War II because of the increased use of petroleum asphalts. Coal tars are also used on airport surfaces because they are very resistant to jet fuels and other petroleum oils. In addition to wearing well, surfaces containing coal tars maintain their skid-resistant qualities over long periods of time.

The importance of coal tar pitches has increased tremendously in recent years because of the growing number of useful chemicals recovered and derived from them. However, the single most important use of coal tar pitch itself in the United States is as a fuel. It is widely used in steel manufacturing as the fuel for open-hearth furnaces. Another important use of pitch is in the manufacture of carbon electrodes, which are widely used in dry cell batteries and in certain metallurgical processes.

Pitch is also used as a foundry core binder and as a binder for the briquetting of coal and coke. Further uses are: in the construction of certain types of roofs; for waterproofing; as a saturant for fiber pipes used to drain water or sewage; as a coating for other types of pipe; as a base in the preparation of various protective coatings; for protection of metals and structures against a marine environment; and to prevent corrosion in underground water, oil, and gas lines.

Creosote, which is one of the heavy tar oils, is of great and long-standing importance as a preservative for wood. It is used to protect objects such as telephone poles, railroad ties, and dock pilings.

While a great number of chemicals may be recovered from the distillation of coal tar, relatively few of these compounds are obtained in large enough quantities to be of real industrial importance. Naphthalene, quinoline, and tar acids, including phenol, cresols, and xylenols, are the most important of the chemicals. Anthracene is also recovered from the coal tar, but in smaller quantities than the other chemicals.

Naphthalene is an intermediate used in the production of dyes, synthetic resins, and other chemicals. It is also used in fungicides, moth repellents, lubricants, and explosives and as a preservative and solvent. Quinoline is used for the production of dyes and niacin and as a preservative and solvent. The coal tar acids are used as insecticides and wood preservatives and in dismicretants. Anthracene is used in the manufacture of dyes.

Mistery. The first reference to tar obtained from coal appeared in a British patent in 1681. However, it was not until 1781 that any commercial uses of coal tar were realized, and even then must recovered coal tar was dumped into the sea. In 1881 the Germans successfully introduced the by-product coking oven for the production of metallurgical grade coke. With the advent of these ovens, it became possible to recover coal tan and pitch in large quantities.

It was gradually realized that important solvents and wood preservatives could be obtained from coal tars and that the residue could be used for highway pavement binders and for specialized custings. Extensive use of the refined coal tars, and especially of coal tar pitches, developed concurrently with the derivation of chemicals and dyes from the distilled fractions of coal tar.

ARNOLD J. HOBERG The Flintkote Company, Whippany, N. J.



Fjords—deep arms of the sea extending far inland—are characteristic of the glaciated coast of Norway.



FAIRCHILD AERIAL SURVEYS

The barrier coast at Lloyd Neck, Long Island, N. Y., is a sand spit separated from the mainland by a lagoon.

COAST, the land bordering an ocean or a sea. Coasts vary considerably according to local geology and geomorphology and the stability of the sea level. Under stable conditions the sea tends to erode the softer rocks and form bays, while the harder rocks remain to form headlands.

FORMATION OF COASTS

Coasts are subject to constant change because of the action of waves, currents, rivers, ice, and winds. They are also subject to occasional change as a result of movements in the earth's crust.

In calm-weather, the sea has little erosional effect on the coast, instead making shoreline deposits of sand and gravel. In stormy weather, great waves hurl sand, gravel, and rocks against the land, causing erosion of the coast.

The material torn loose from the land by the sea is graded according to size. Gravel and sand remain close to shore, forming beaches, spits, and bars, while the fine material is swept out to sea. The speed with which the sea erodes the coast depends on the frequency of powerful onshore winds, the resulting waves, and the resistance of the coastal rocks.

Roberts + Caserio. Basic Principles of Organic Chemistry NY; W.A. Benjamin, Inc., 1965

Sec 3-5 | Estimation of Heats of Combustion

75

whether an alkane is branched or not can be readily ascertained by inspection of the ratio of the integral of the CH₃ resonances centered on 0.9 ppm versus the integral of the CH₂ resonances centered on 1.25 ppm (see Figure 3-4).

Mass spectrometry is excellent for the analysis of alkane mixtures and is widely used for this purpose.

CHEMICAL REACTIONS OF ALKANES

3-4 COMBUSTION OF ALKANES

As a class, alkanes are singularly unreactive. Hence the name saturated hydrocarbon or "paraffin," which literally means "not enough affinity" [L. par(um), not enough, + affins, affinity], arises because their chemical "affinity" for most common reagents may be regarded as "saturated" or satisfied. Thus none of the C-H or C-C bonds in a typical saturated hydrocarbon, such as ethane, are attacked at ordinary temperatures by a strong acid such as sulfuric acid or by an oxidizing agent such as bromine (in the dark), oxygen, or potassium permanganate. Ethane is similarly stable under ordinary conditions to reducing agents such as hydrogen in the presence of such catalysts as platinum, palladium, or nickel.

However, all saturated hydrocarbons are attacked by oxygen at elevated temperatures and, if oxygen is in excess, complete combustion occurs to carbon dioxide and water. Vast quantities of hydrocarbons from petroleum are utilized as fuels for the production of heat and power by combustion. Although petroleums differ in omposition with their source, a representative petroleum on distillation yields the following fractions: (a) Gas fraction, boiling point up to 40° C, contains normal and branched alkanes from C1 to C5. Natural gas is mainly methane and ethane while "bottled" gas (liquefied petroleum gas) is mainly propane and butane. (b) Gasoline, boiling point from 40 to 180° C, contains hydrocarbons from C₆ to C₁₀. Over 100 compounds have been identified in gasoline, and these include normal and branched alkanes, cycloalkanes, and alkylbenzenes (arenes). The branched alkanes make better gasoline than their straight-chain isomers because they have much higher antiknock ratings. (c) Kerosine, boiling point 180 to 230° C, contains hydrocarbons from C11 to C12. Much of this is utilized as jet engine fuels and much is "cracked" to simpler alkanes (and alkenes). (d) Light gas oil, boiling point 230 to 305° C, C13 to C17, is utilized as diesel and furnace fuels. (e) Heavy gas oil and light lubricating distillate, boiling point 305 to 405° C, C₁₈ to C₂₅. (f) Lubricants, boiling point 405 to 515° C, C₂₆ to C₃₈, familiarly encountered as paraffin wax and petroleum jelly (Vaseline). The distillation residues are better known as asphalts.

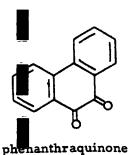
3-5 ESTIMATION OF HEATS OF COMBUSTION. BOND ENERGIES

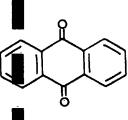
The combustion of alkanes obtained from petroleum is a major source of heat and power. Consequently, it is of practical interest to be able to estimate the amount of heat liberated in the combustion of different kinds of hydrocarbons. This can be done quite simply with the aid of tables of bond energies—and the method is See "Hydrocarbons in Petroleum" by F. D. Rossini, J. Chem. Ed., 37, 554 (1960).

Sec. 22-10 | Sources and Uses of Aromatic Hydrocarbons

itic Substitution | Chap. 22

renzene or naphthanthraquinone, and





nthraquinone

base onsiderably more stable

.

Bentane itself gives glyoxal.

The double-bond character of the 9,10-bond in phenanthrene is particularly evident in ozonization. This bond is attacked preferentially and leads to the formation of 2,2'-diphenaldehyde when the ozonide is reduced with iodide ion.

2,2'-diphenaldehyde

EXERCISE 22-42 What products would you expect to be formed in the ozonization of the following substances (consider carefully which bonds are likely to be most reactive)?

a. o-xylene

b. naphthalene

22-10 SOURCES AND USES OF AROMATIC HYDROCARBONS

Benzene and many of its derivatives are manufactured on a very large scale for use in high-octane gasolines, and in the production of polymers, insecticides, detergents, dyes, and other miscellaneous chemicals. Prior to World War II, coal was the only important source of aromatic hydrocarbons, but, during the war and thereafter, the demand for benzene, toluene, and the xylenes rose so sharply that other sources had to be found. Today, about 70 per cent of the benzene and almost all of the toluene and the xylenes produced in the United States are derived from petroleum.

A. Aromatics from Coal

When coal is heated at high temperatures in the absence of air, it carbonizes to coke and gives off a gaseous mixture of compounds, some of which condense to a black viscous oil (coal tar), others to an aqueous condensate called ammoniacal liquors, and some remain gaseous (coal gas). Crude coal gas contains small but valuable amounts of benzene, toluene, and the xylenes mixed with many other compounds including cyclopentadiene, thiophene, and naphthalene. The arenes are recovered as light oil by a process known as "scrubbing" and are then separated by fractional distillation.

Very little is wasted in the coking of coal. Coal tar is the source of an amazing number of compounds, some of which are listed in Table 22-9; they are mainly aromatic and in-

Principal Compounds Obtained from Coal Tarab Table 22-9

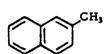
Hydrocarbonse



naphthalene



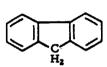
l-methyl-naphthalene



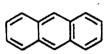
2-methyl-naphthalene



acenaphthene



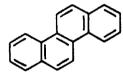
fluorene



anthracene



phenanthrene

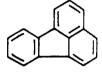


chrysene

į



pyrene



fluoranthene

Nitrogen compounds



pyridine

α-picoline

β-picoline

y-picoline

quinoline

carbazole

acridine

- Trincipal Con	iipoonas Oblainea from	Continued)			
Oxygen compounds					
OH OH	СН,	CH3	CH ₃		
phenol	<i>o</i> -cresol	<i>m</i> =cresol	P-cresol		
CH ₃ OH		OH	ОН		
p-xylenol (and other xylenols)	l÷nap	hthol	2-naphthol		

- ^aCompiled from "Chemistry of Coal Utilization," National Research Council Committee, H. H. Lowry (ed.), Wiley, 1945.
- None of the compounds listed are very abundant in coal tar. Even naphthalene, which is present in the largest amount, constitutes only 10 to 11 per cent of coal tar.
- ^c Benzene, toluene, and the xylenes are present in very small amount (0.3 per cent total) in coal tar.

clude oxygen and nitrogen compounds as well as hydrocarbons. Coal gas (freed of light oil) is used as a fuel and the ammoniacal liquors are a source of ammonia. The residual coke is used both as a fuel and as a source of carbon for the production of steel from iron oxides. In fact, one of the reasons that new sources of arenes had to be found was the undesirable dependence of the supply of arenes on the demand for coke by the steel industry.

It is also possible to make a variety of aromatic hydrocarbons by hydrogenation of powdered coal at high temperature and pressure. Such processes do not appear to be economically feasible at the present time.

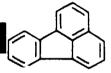
B. Aromatics from Petroleum

Although petroleums from some locations do possess fairly substantial amounts of aromatic hydrocarbons, they are not a principal source of supply for such compounds. Rather, aromatics are synthesized from the aliphatic components of the C_6 — C_{10} gasoline fraction from petroleum refining by a process that is referred to in the petrochemical industry as catalytic reforming. This involves passing the hydrocarbon vapors mixed with excess hydrogen over a catalyst, usually platinum on alumina, at elevated temperatures (900°C) and pressures (200 to 500 psi). Several remarkable transformations take place. For example: Cyclohexane and methylcyclohexane are dehydrogenated to benzene and toluene, respectively;



acenaphthene

phenanthrene



fluoranthene



γ-picoline

acridine

dimethylcyclopentanes are isomerized to methylcyclohexane, which is then dehydrogenated to toluene;

$$H_3C$$
 CH_3
 GH_3
 and open-chain alkanes are cyclized to cycloalkanes, which then lose hydrogen to give arenes.

n-heptane

In spite of the fact that hydrogen is a product of reforming, excess hydrogen is necessary to reduce formation of carbon deposits on the catalyst. As a further precaution against poisoning the catalyst, the reactants are freed from nitrogen and sulfur contaminants by first heating them in the presence of hydrogen and a cobalt-molybdenum catalyst, then fractionating the hydrocarbons from the ammonia and hydrogen sulfide thereby formed.

Much of the aromatic product obtained by catalytic reforming is blended with other fractions from petroleum refining to give high-octane gasoline. The rest is separated into its component hydrocarbons, which are then utilized by the chemical industry for the production of chemicals derived from benzene, toluene, and the xylenes (see Table 22-10).

The ratio of benzene, toluene, and xylenes obtained from petroleum is roughly 1:4:5, which contrasts with the ratio of 16:3:1 obtained from coal. Since there is currently more demand for benzene than for toluene, much of the toluene obtained from petroleum is converted to benzene by a process called hydrodealkylation. This involves heating a mixture of toluene with excess hydrogen to fairly high temperatures in the presence of a cobalt-molybdenum catalyst.

Naphthalene is obtained primarily from coal tar, but increasing amounts are being produced from petroleum. The residues from catalytic reforming of gasoline are rich in methylnaphthalenes, and these can be hydrodealkylated to naphthalene in the same way that toluene is converted to benzene.

U. S. DEPARTMENT OF COMMERCE WEATHER BUREAU

CLIMATOGRAPHY OF THE UNITED STATES NO. 60 - 30

CLIMATES OF THE STATES

NEW YORK



WASHINGTON, D. C.

FEBRUARY 1960

Climate of New York

Ernest C. Johnson, Weather Bureau State Climatologist

The climate of New York State is broadly representative of the moist continental type area which liankets the northeastern United States, but its diversity is not usually encountered within such a mall area. Differences in latitude, character of typography, and bodies of water have pronounced effects on the climate of local areas. The geographical position of the State and the usual curse of air masses governed by the atmospheric circulation of the region provide the general climatic controls.

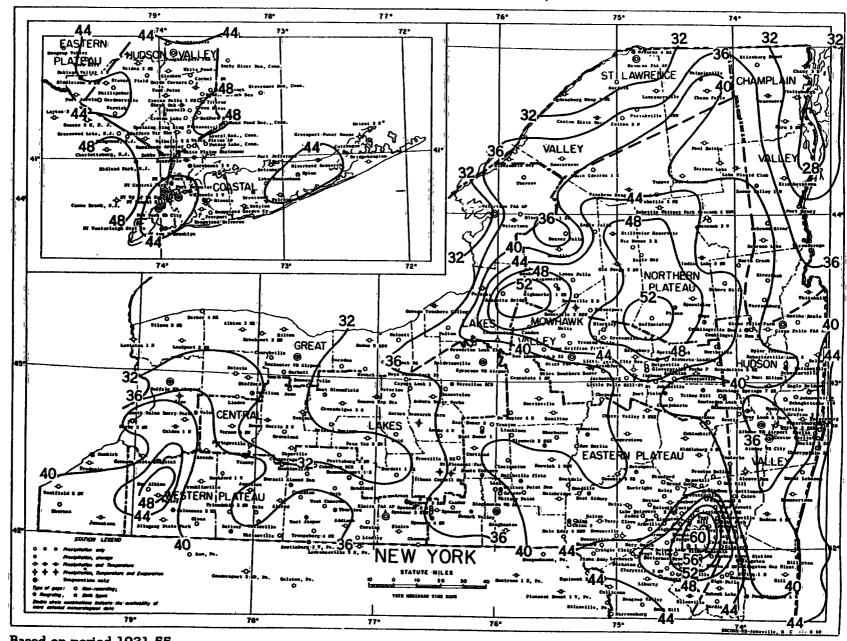
New York State contains 49,576 square miles exclusive of the boundary water areas of Long island Sound. New York Harbor, Lake Ontario, and lake Erie. The major portion of its area lies approximately between latitudes 42°N. and 45°N.; longitudes 73°30'W. and 79°45'W. However, in the extreme southeast a triangular portion extends southward to latitude 41°N., while Long island lies eastward between latitudes 41°00'N. and 40°35'N. and longitudes 72°W. and 74°W.

The principal highlands of the State may be divided into two general regions, namely, the Idirondack section in the northeast and the Appalachian plateau in the southern portion west of the Hudson Valley. A subdivision of the Appalachian highlands is produced by the deep channel of Seneca Lake, extending from the plains bordering Lake Ontario southward to the Susquehanna Valley. Thus are formed the areas commonly called

the Eastern and Western Plateaus; the former extending from the central lakes to the Hudson Valley and the latter westward from the central lakes to the depression of Lake Erie. The Eastern Plateau includes the Catskill Mountains, which are the northeastern New York terminus of the Allegheny Range of the Appalachian Mountain system. In southeastern New York is a minor highland region cut through by the Hudson River. This includes the Highlands, the Palisades, and the Taconic Mountains. Another minor highland known as Tug Hill lies just west of the Adirondacks and the Black River, and includes a large part of Lewis County.

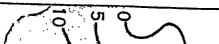
Along the eastern border of the State is a long. narrow lowland, which is occupied by Lake Champlain, Lake George, and the middle and lower portions of the Hudson Valley. The St. Lawrence River, Lake Ontario, and Lake Erie lie in another lowland region on the north and western boundaries. This region is widest south of the eastern end of Lake Ontario, narrowing to a width of only 4 or 5 miles in the western end of the State. Connecting the Hudson-Champlain Valley with the lake plains is a third lowland. This is a deep transversal cut which forms the valley of the Mohawk River and Lake Oneida, and thus divides the main plateau area into the distinct masses of the Appalachian and Adirondack systems. A fourth lowland region is Long Island which is a part of the Atlantic

Mean Annual Precipitation, Inches



Based on period 1931-55

Isolines are drawn through points of approximately equal value. Caution should be used in interpolating on these maps, particulary in mountainous areas.



NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION DIVISION OF SOLID AND HAZARDOUS WASTE INACTIVE HAZARDOUS WASTE DISPOSAL SITE REPORT
CLASSIFICATION CODE: 2a REGIÓN: 6 SITE CODE:
NAME OF SITE: New York Emulsions Tar Products STREET ADDRESS: Washington Street TOWN/CITY: COUNTY: ZIF: Operida: 13501
SITE TYPE: Open DumpStructure-XLagoonLandfillTreatment Pond ESTIMATED SIZE:
SITE OWNER/OPERATOR INFORMATION: CURRENT OWNER NAME: CURRENT OWNER ADDRESS: Washington St., Which OWNER(S) DURING USE: OPERATOR DURING USE: Same OPERATOR ADDRESS: PERIOD ASSOCIATED WITH HAZARDOUS WASTE: From To
This is a distribution center for tor emulsions. The site consists of numerous tanks and associated piping.
A remedial investigation completed by Niagara Mohowll Power Corp. indicates that this site may be a source of naphthalene and benzene confamination in
the groundwater. This facility is no longer operating.
HAZARDOUS WASTE DISPOSED: Confirmed Suspected
na phthalene benzene

ANALYTICAL DATA AVAILABLE:	SITE CODE:
AirSurface WaterGroundwaterXS	SoilSediment- None-
CONTRAVENTION OF STANDARDS:	
Groundwater-XIrinking WaterSurfac	ce WaterAir
LEGAL ACTION:	
TYPE:State—X Federal— STATUS:In Progress—X Completed—	
REMEDIAL ACTION:	
Proposed-XUnder DesignIn Progress NATURE OF ACTION:	Completed
GEOTECHNICAL INFORMATION: SOIL TYPE: Fluvial deports, glaciólacustria e GROUNDWATER DEPTH: 1-9 \$+	deposits as glucial till.
ASSESSMENT OF ENVIRONMENTAL PROBLEMS:	
: groundwater contamination	
• •	
•	
ASSESSMENT OF HEALTH PROBLEMS:	
•	••
•	
•	•
PERSON(S) COMPLETING THIS FORM:	
NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION	NEW YORK STATE DEPARTMENT OF HEALTH
NAME: Darrell Sweredoski TITLE: Senior Sanitary Eng.	NAME: TITLE:
NAME: TITLE:	NAME: TITLE:
DATE: Nov. 21, 1986	DATE:

DRAFT

SUBJECT TO REVISION NOT FOR EXTERNAL RELEASE New York State
Department of Environmental Conservation
50 Wolf Road
Albany, New York 12233

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION SUMMARY REPORT

New York Emulsions Tar Products Site Name Washington Street, Utica, Oneida C. Address/County	DEC Site ID Number (Registry Sites Only)
SITE DESCRIPTION distribution center for ter emulsion vicinity of the site contomerate and naphthalene	Date of Visit ions; groundwater in led with benzene
Phase I # II	
PRIORITY FOR FURTHER ACTION High	Medium X Low
ADD TO REGISTRY X Yes No	2a Suggested Classification
JUSTIFICATION (yes or no) RI completed by Miagara Mohaus groundwater contaminates may be site	
Prepared by: <u>Jamell Sweredak</u> : Dar	te: Nov 21, 1986



Preliminary Assessment

	POTENTIAL HA	ZARDOUS W	ASTE SITE	L.	I. IDENTIFICATION	
SEPA		RY ASSESS		į,	1 STATE 02 SITE NUMBER	
	PART 1 - SITE INFOR	MATION AND	ASSESSMEN	т 4	(XXXX)	
II. SITE NAME AND LOCATION				J	· · · · · · · · · · · · · · · · · · ·	
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III. RESPONSIBLE PARTIES					and almost the second	
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Utica			13501	()		
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D9 CITY		10 STATE 11	ZIP CODE	12 TELEPHONE NI	IMBER	-
				()	-	
13 TYPE OF OWNERSHIP (Check one)			 			
A PRIVATE C B FEDI	ERAL (Agency name)		C. STATE	DD.COUNTY	☐ E. MUNICIPAL	
□ F. OTHĒR	(Specify)		E G UNKNOW			
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IV. CHARACTERIZATION OF POTEN	ITIAL HAZARD	, , , , , , , , , , , , , , , , , , ,				
01 ON SITE INSPECTION	BY (Check all that apply)	50.00.00.00				
THE DATE / /	□ A. EPA □ B E. LOCAL HEALTH	EPA CONTRACT OFFICIAL DEF	OTHER	STATE D	D. OTHER CONTRACTOR	
2.10	CONTRACTOR NAME(Si		(\$4	ecry)	
02 SITE STATUS (Check one)	03 YEARS OF C					
□ A. ACTIVE SE B INACTIVE □	C UNKNOWN		_1	5<	UNKNOWN	
04 DESCRIPTION OF SUBSTANCES POSSIBLY	PRESENT, KNOWN OR ALLEGED	BEGINNING YEAR	ENDING YEA	<u> </u>		
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	<u> </u>					
05 DESCRIPTION OF POTENTIAL HAZARD TO	ENVIRONMENT AND OR POPULATION				* (<u>L. 5,9</u> *)	
V. PRIORITY ASSESSMENT						
01 PRIORITY FOR INSPECTION (Greek one # App		Information and Part 3		us Conditions and Incide	nts)	***************************************
A HIGH (Properties required premptly)		n time avedeble basis)	D NONE (No further a	chan neodod complete	current disposition form)	
	VI. INFORMATION AVAILABLE FROM					
01 CONTACT	02 OF (Agency O		. /	/	03 TELEPHONE NU	JMBER
Darrell Swered	oski NYS	DEC.	Wasker	town	13/57 785-	252
04 PERSON RESPONSIBLE FOR ASSESSMEN	T 05 AGENCY	06 ORGANIZ	ATION	07 TELEPHONE N	UMBER OB DATE	
Same				()	MONTH DAY	FAR .
EPA FORM 2070-12 (7-81)						*F.F.



ŞEPA

POTENTIAL HAZARDOUS WASTE SITE PRELIMINARY ASSESSMENT PART 2. WASTE INFORMATION

I. IDENTIFICATION
O1 STATE G2 SITE NUMBER
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	, ·		PART 2 - WAST	E INFORMATION	l .		
II WASTES	TATES, QUANTITIES, AN	ND CHARACTER	İSTICS		3,5-50,55		
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D OTHER	Scocii.	NO OF DRUMS				M NOT AP	PLICABLE
III. WASTE T	YPE						•
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OFM	OILY WASTE				**************************************		•
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PSD	PESTICIDES					- 1-14	/
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, юс	INORGANIC CHEMIC	CALS			_		<u> </u>
ACD	ACIDS						
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MÈS	HEAVY METALS					der to the state of the state of	· · · · · · · · · · · · · · · · · · ·
	OUS SUBSTANCES 500 AS	Spendia for most frequent	riy crea CAS humbers				
C1 CATEGORY	02 SUBSTANCE N	IAME	03 CAS NUMBER	04 STORAGE DISF	POSAL METHOD	05 CONCENTRATION	DE MEASURE OF CONCENTRATION
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	S OF INFORMATION Const				XXXXXXXXX	XXXXXXXXXX	XXXXXXXX.
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POTENTIAL HAZARDOUS WASTE SITE		I. IDENTIFICATION	
	MINARY ASSESSMENT	CT STATE D2 SITE NUMBER	
PART 3 - DESCRIPTION OF	HAZARDOUS CONDITIONS AND INCIDENTS	XXXX	
II. HAZARDOUS CONDITIONS AND INCIDENTS			
01 XA GROUNDWATER CONTAMINATION		POTENTIAL XALLEGED	
03 POPULATION POTENTIALLY AFFECTED	04 NARRATIVE DESCRIPTION	_	
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01 VR SUBSACE WATER CONTAMINATION	02 - OBSERVED (DATE	POTENTIAL _ ALLEGED	
01 XB SURFACE WATER CONTAMINATION 03 POPULATION POTENTIALLY AFFECTED	04 NARRATIVE DESCRIPTION		
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03 POPULATION POTENTIALLY AFFECTED	02 OBSERVED (DATE)	POTENTIAL _ ALLEGED	
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01 X F CONTAMINATION OF SOIL	02 T OBSERVED (DATE) =	POTENTIAL ALLEGED	
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Strong potential	CX1575		
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POTENTIAL HAZARDOUS WASTE SITE PRELIMINARY ASSESSMENT PART 3 - DESCRIPTION OF HAZARDOUS CONDITIONS AND INCIDENTS		I. IDENTIF	
		XXXX	? SITE NUMBER
II. HAZARDOUS CONDITIONS AND INCIDENTS COMPAND			
01 I J DAMAGE TO FLORA 04 NARRATIVE DESCRIPTION Unknown		POTENTIAL	C ALLEGED
01 I K DAMAGE TO FAUNA 04 NARRATIVE DESCRIPTION INClude name/31 of speciess Len Une	· -	POTENTIAL	C ALLEGED .
01 I L CONTAMENATION OF FOOD CHAIN 04 NARRATIVE DESCRIPTION Len known	· · · · · · · · · · · · · · · · · · ·	POTENTIAL	C ALLEGEÓ
01 & M UNSTABLE CONTAINMENT OF WASTES (Speed nation) standing bounds making drums. 03 POPULATION POTENTIALLY AFFECTED Lugh puter trail for	02 = OBSERVED (DATE) 04 NARRATIVE DESCRIPTION Confaminade inigration	(POTENTIAL	C ALLEGED .
01 = N DAMAGE TO OFFSITE PROPERTY 04 NARRATIVE DESCRIPTION CENTROL	n	POTENTIAL	S ALLEGED
01 TO CONTAMINATION OF SEWERS STORM DRAINS WATPS 04 MARRATIVE DESCRIPTION Migh patential due	to infiltration of	POTENTIAL 9	E ALLEGED
01 T P ILLEGAL/UNAUTHORIZED DUMPING 04 NARRATIVE DESCRIPTION	02 TOBSERVED (DATE)	POTENTIAL	Ë ALLEGED
05 DESCRIPTION OF ANY OTHER KNOWN, POTENTIAL, OR ALLEG	ED HAZARDS		
HI. TOTAL POPULATION POTENTIALLY AFFECTED:			
IV. COMMENTS			
in the area which ing gource of groundwater	licetes that this is	ted a	n KI y be a
V. SOUNCES OF INFORMATION (Cité specific references e g. state lites sample analysis reports)			
1/2 gara Molwell 1/arbor Point Prop	RI March 1985 certy	,	

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New York Emulsions Tar Producte Dir Suit-Kote Corp Washington St., Utica Main Office - Cortland - Suit-Kote Corp. Herfer, Secrest, & Emery Alborray I have 700 illidtown Tower, Rochester R. Clinton Enery



New York State Department of Environmental Conservation

MEMORANDUM

TO:

Walt Demick

FROM:

Darrell Sweredoski, Region 6

SUBJECT: NEW YORK EMULSIONS TAR PRODUCTS, UTICA - #633031

DATE:

July 14, 1987

On this date I was contacted by Mr. Mark Kepple from Environmental Oils. Mr. Kepple stated that he was representing a client involved with the above site. He was not at liberty to divulge the client's name.

Mr. Kepple asked me about the status of the New York Emulsion site. \hat{I} explained to him that it was not currently on the registry, but I expected it to appear in the July registry. I also informed him that the site was scheduled for a Phase I investigation.

I went on to describe to Mr. Kepple our concerns about the site. The Ni-Mo Power Corporation had done a remedial investigation on adjacent property and concluded that groundwater contamination in the form of naphthalene and benzene were eminating from the New York Emulsions site. I also informed him that since monitoring wells were not actually placed on the New York Emulsion site, that the site was only listed as a 2A suspect site.

Mr. Kepple suggested that his client would be interested in a meeting to discuss the issues. I told him that any such meetings would involve our Central Office technical staff and possibly our legal staff. Mr. Kepple said he would discuss the situation with his client and get back in touch with us.

Darrell M. Sweredoski, P.E.

Sr. Sanitary Engineer

Division Solid/Hazardous Waste

DMS: kw

NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION DIVISION OF HAZARDOUS WASTE REMEDIATION INACTIVE HAZARDOUS WASTE DISPOSAL REPORT

CLASSIFICATION CODE: 2

REGION: 6

SITE CODE: 633021 EPA ID: NYD980664411

NAME OF SITE: Niagara Mohawk Harbor Pt. Property

STREET ADDRESS: Washington Street

COUNTY:

ZIP:

TOWN/CITY: Utica

Oneida

13502

SITE TYPE: Open Dump-X Structure- Lagoon- Landfill- Treatment Pond-

ESTIMATED SIZE: 64

SITE OWNER/OPERATOR INFORMATION:

CURRENT OWNER NAME....: Niagara Mohawk Power Corp.

CURRENT OWNER ADDRESS .: 300 Erie Blvd. West, Syracuse, NY

OWNER(S) DURING USE...: Utica Gas & Electric

OPERATOR DURING USE ...: Niagara Mohawk Power Corp.

OPERATOR ADDRESS.....: 300 Erie Blvd. West, Syracuse, NY

To 1950's PERIOD ASSOCIATED WITH HAZARDOUS WASTE: From 1902

SITE DESCRIPTION:

Site of former Water Gas and Coal Gasification Plants operated by Utica Gas and Electric. Gas was passed through purifier boxes containing wood shavings and iron oxides to remove cyanides. Mixture from purifier box and ash spread over various areas of site. Spillage from product storage areas and coke plant operation. Neighboring operations include an Asphalt Emulsions Plant and Chemical Company. Coal tars spilled on site. Two additional tank farms nearby. A portion of an old municipal landfill is located on this site. DEE is planning to begin negotiations for an RI/FS in June of 1989.

Suspected-Confirmed-X HAZARDOUS WASTE DISPOSED: QUANTITY (units) TYPE 24,800 cu yds. cyanides (spent mixture) 60,000 cu yds. coal tars 15,200 cu yds. tar and oil

SITE CODE: 633021

ANALYTICAL DATA AVAILABLE:

Surface Water-X Groundwater-X Soil-X Sediment-X

CONTRAVENTION OF STANDARDS:

Drinking Water- Surface Water-Groundwater-X

Air-

LEGAL ACTION:

Federal-State- X TYPE..: Consent Order Negotiation in Progress- X Order Signed-

REMEDIAL ACTION:

Under design- In Progress-Completed-Proposed-X

NATURE OF ACTION: RI-FS

GEOTECHNICAL INFORMATION:

SOIL TYPE: waste fill, clays, sand gravel, silty sands & clays

GROUNDWATER DEPTH: 2-10 feet

ASSESSMENT OF ENVIRONMENTAL PROBLEMS:

Niagara Mohawk has completed initial studies of land and Mohawk River. Releases of cyanide, benzene and polynuclear aromatics to shallow groundwaters and Mohawk River confirmed. Deep aquifer studies on-going.

Heavy soil contamination on site. Site was fenced 11/84 by Niagara Mohawk.

ASSESSMENT OF HEALTH PROBLEMS:

Direct contact concerns caused by heavy onsite soil contamination have been addressed by the erection of a fence at DOH request (11/84). Therefore, no exposure to soil is likely, unless movement or use of contaminated soil occurs. This may be a consideration if plans to develop the area for commercial/recreational purposes proceed. Shallow groundwater contamination and potential deep aquifer contamination discharge to the Mohawk River, posing potential recreational hazards. The area is on public water. However, DOH will reinspect the area in summer, 1989 to reconfirm that no private wells in the area continue to be used for drinking water, unless they are sampled.

NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION DIVISION OF HAZARDOUS WASTE REMEDIATION INACTIVE HAZARDOUS WASTE DISPOSAL REPORT

CLASSIFICATION CODE: 2a

REGION: 6

SITE CODE: 633030

EPA ID:

NAME OF SITE: Monarch Chemical Company Inc.

STREET ADDRESS: 37 Meadow Street

TOWN/CITY:

COUNTY: Oneida

ZIP: 13503

SITE TYPE: Open Dump-X Structure- Lagoon- Landfill- Treatment Pond-

ESTIMATED SIZE: 6

SITE OWNER/OPERATOR INFORMATION:

CURRENT OWNER NAME...: Monarch Chemical Co.Inc. CURRENT OWNER ADDRESS .: 37 Meadow St., Utica, NY

OWNER(S) DURING USE...: Same as above

OPERATOR DURING USE ...: Monarch Chemical Company, Inc.

OPERATOR ADDRESS.....: 37 Meadow Street, Utica, NY

PERIOD ASSOCIATED WITH HAZARDOUS WASTE: From 1966 To present

SITE DESCRIPTION:

Utica

Chemical distribution/manufacturing plant of sodium hydroxide and sodium hypochlorite. The plant uses various acids in their production process. They also receive and distribute other assorted chemicals. Site is located in an industrial area adjacent to the Mohawk River and Utica Harbor. Groundwater analysis show contamination from dichloroethanes, trichloroethanes and dichloroethylenes. A Phase I study is currently underway, and a Phase II study is planned.

HAZARDOUS WASTE DISPOSED: Confirmed-X

Suspected-QUANTITY (units)

dichloroethanes dichloroethylenes

trichloroethane

unknown

SITE CODE: 633030

ALYTICAL DATA AVAILABLE:

Air- Surface Water- Groundwater-X Soil- Sediment-

ONTRAVENTION OF STANDARDS:

Groundwater-X Drinking Water- Surface Water- Air-

I GAL ACTION:

TYPE..: SEATUS: State-

Federal-

Negotiation in Progress- Order Signed-

REMEDIAL ACTION:

Hoposed- Under design-

In Progress-

Completed-

NATURE OF ACTION:

TOTECHNICAL INFORMATION:

SIL TYPE:

GROUNDWATER DEPTH: 5 feet

A SESSMENT OF ENVIRONMENTAL PROBLEMS:

Groundwater beneath the site is contaminated with various solvents.

ASSESSMENT OF HEALTH PROBLEMS:

NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION DIVISION OF HAZARDOUS WASTE REMEDIATION INACTIVE HAZARDOUS WASTE DISPOSAL REPORT

SITE CODE: 633032 CLASSIFICATION CODE: 2a REGION: 6

EPA ID: NYD986866010

NAME OF SITE : Mohawk Valley Oil Inc.

STREET ADDRESS: Washington Street

ZIP: COUNTY: TOWN/CITY: 13501 Oneida Utica

SITE TYPE: Open Dump- Structure-X Lagoon- Landfill- Treatment Pond-

Acres ESTIMATED SIZE: 3.5

SITE OWNER/OPERATOR INFORMATION:

CURRENT OWNER NAME...: ** Multi - Owner Site **
CURRENT OWNER ADDRESS . * * * * *

CURRENT OWNER ADDRESS .:

OWNER(S) DURING USE...: Mohawk Valley Oil Inc. OPERATOR DURING USE ...: Mohawk Valley Oil Co. OPERATOR ADDRESS.....: Route 49, Marcy, NY

PERIOD ASSOCIATED WITH HAZARDOUS WASTE: From To

SITE DESCRIPTION:

This was a tank storage area used to store fuel. There are 2 separate sections of land the comprises this site. The active portion of the property is the Mohawk Valley Oil Terminal. The other section is the former Niagara Flats Terminal which was originally owned by the Niagara-Mohawk Power Co. This property was purchased by the Mohawk Valley Oil Co. in 1961 and sold to the Interstate Harbor Development in September of 1987.

A remedial investigation completed by NI-MO Power Co. indicates that this site may be the source of napthalene and benzene contamination in the groundwater. The Mohawk Valley oil terminal tanks have been removed and the Niagara Flats property has been filled in with construct ion and demolition material. A Phase I study is underway and a Phase II

is planned for 1989.

Suspected-X HAZARDOUS WASTE DISPOSED: Confirmed-QUANTITY (units)

Unknown Benzene

Napthalene

SITE CODE: 633032

ANALYTICAL DATA AVAILABLE:

ir- Surface Water- Groundwater-X Soil- Sediment-

CONTRAVENTION OF STANDARDS:

Air-Froundwater-X Drinking Water- Surface Water-

LEGAL ACTION:

Federal-TYPE..: State- Federal-STATUS: Negotiation in Progress- Order Signed-State-

REMEDIAL ACTION:

Proposed- Under design- In Progress- Completed-

NATURE OF ACTION:

GEOTECHNICAL INFORMATION:

SOIL TYPE: Glacial Lacustrine deposits and Glacial Till

FROUNDWATER DEPTH: 1-9 Feet

ASSESSMENT OF ENVIRONMENTAL PROBLEMS:

Contamination of groundwater

ASSESSMENT OF HEALTH PROBLEMS:

6.2 <u>APPENDIX B - REVISED NYSDEC INACTIVE HAZARDOUS WASTE</u>
<u>DISPOSAL SITE REPORT</u>

(47-15-11 (10/83)

NEW YORK STATE DÉPARTMENT OF ENVIRONMENTAL CONSERVATION DIVISION OF SOLID AND HAZARDOUS WASTE INACTIVÉ HAZARDOUS WASTE DISPOSAL SITE REPORT

EGION: 6

LAGOON
related products, on 2.96 acres as source of also have
OUNDS, DRUMS, TONS, GALLONS)

TIME PERIOD SITE WAS USED FOR HAZARD	
	5 TOapprox. , 19 83
· ·	ers Products Co.; Suit-Kote Corp.
SITE OPERATOR DURING PERIOD OF USE:	
	: 1911 Lorings Crossing Road, Cortland, N.Y. 1304
• • • • • • • • • • • • • • • • • • •	SURFACE WATER X GROUNDWATER X SEDIMENT X NONE
CONTRAVENTION OF STANDARDS: GROUN SURFA	DWATER DRINKING WATER CE WATER AIR
SOIL TYPE: Unconsolidated fill, sil DEPTH TO GROUNDWATER TABLE: 10-15 f	
DEPTH TO GROUNDMATER TABLE: 10-13 1	
LEGAL ACTION: TYPE:	STATE FEDERAL
STATUS: IN PROGRESS	COMPLETED
REMEDIAL ACTION: PROPOSED	UNDER DESIGN
IN PROGRESS	COMPLETED
NATURE OF ACTION:	
naphthalene and benzene contamination	a Mohawk, owner of adjacent property, indicates on of groundwater from this site. Other r sediments may be affected. Water column
ASSESSMENT OF HEALTH PROBLEMS:	
Some risk of direct contact. Fairly route. 120 potable water wells known 15 miles away.	y low risk by groundwater or surface water wn within three mile radius. Surface water intakes
PERSON(S) COMPLETING THIS FORM:	
NEW YORK STATE DÉPARTMENT OF ENVIRONMENTAL CONSERVATION	NEW YORK STATE DEPARTMENT OF HEALTH
NAME Kevin H. Siepel	NAME
TITLE Scientist-URS Corp.	TITLE
NAME	NAME
TITLE	TITLE
DATE: November 11, 1987	DATE: